COMPOSITIONS OF ORTHOGONAL LEUCYL-tRNA AND AMINOACYL-tRNA SYNTHETASE PAIRS AND USES THEREOF

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001]   This application claims priority to and benefit of Provisional Patent Application USSN 60/485,451, filed July 7, 2003; and to Provisional Patent Application USSN 60/488,215, filed July 16, 2003, the disclosures of which are incorporated herein by reference in their entirety for all purposes.

STATEMENT AS TO RIGHTS TO INVENTIONS MADE UNDER FEDERALLY SPONSORED RESEARCH AND DEVELOPMENT

[0002]   This invention was made with government support under Grant No. GM 62159 from the National Institutes of Health. The government may have certain rights to this invention.

FIELD OF THE INVENTION

[0003]   The invention pertains to the field of translation biochemistry. The invention relates to methods for producing and compositions of orthogonal leucyl tRNAs, orthogonal leucyl aminoacyl-tRNA synthetases and pairs thereof. The invention also relates to methods of producing proteins in cells using such pairs and related compositions.

BACKGROUND OF THE INVENTION

[0004]   The genetic code of every known organism, from bacteria to humans, encodes the same twenty common amino acids. Different combinations of the same twenty natural amino acids form proteins that carry out virtually all the complex processes of life, from photosynthesis to signal transduction and the immune response. In order to study and modify protein structure and function, scientists have attempted to manipulate both the genetic code and the amino acid sequence of protein. However, it has been difficult to remove the constraints imposed by the genetic code that limit proteins to twenty genetically encoded standard building blocks (with the rare exception of selenocysteine (*see, e.g.,* A. Bock et al., (1991), <u>Molecular Microbiology</u> 5:515-20) and pyrrolysine (*see, e.g.,* G. Srinivasan, et al., (2002), <u>Science</u> 296:1459-62).

[0005]   Some progress has been made to remove these constraints, although this progress has been limited and the ability to rationally control protein structure and function is still in its infancy. For example, chemists have developed methods and strategies to synthesize and manipulate the structures of small molecules (*see, e.g.,* E. J. Corey, & X.-M. Cheng, The Logic of Chemical Synthesis (Wiley-Interscience, New York, 1995)). Total synthesis (*see, e.g.,* B. Merrifield, (1986), Science 232:341-7 (1986)), and semi-synthetic methodologies (*see, e.g.,* D. Y. Jackson et al., (1994) Science 266:243-7; and, P. E. Dawson, & S. B. Kent, (2000), Annual Review of Biochemistry 69:923-60), have made it possible to synthesize peptides and small proteins, but these methodologies have limited utility with proteins over 10 kilo Daltons (kDa). Mutagenesis methods, though powerful, are restricted to a limited number of structural changes. In a number of cases, it has been possible to competitively incorporate close structural analogues of common amino acids throughout proteins. *See, e.g.,* R. Furter, (1998), Protein Science 7:419-26; K. Kirshenbaum, et al., (2002), ChemBioChem 3:235-7; and, V. Doring et al., (2001), Science 292:501-4.

[0006]   Early work demonstrated that the translational machinery of *E. coli* would accommodate amino acids similar in structure to the common twenty. *See,* Hortin, G., and Boime, I. (1983) Methods Enzymol. 96:777-784. This work was further extended by relaxing the specificity of endogenous *E. coli* synthetases so that they activate unnatural amino acids as well as their cognate natural amino acid. Moreover, it was shown that mutations in editing domains could also be used to extend the substrate scope of the endogenous synthetase. *See,* Doring, V., et al., (2001) Science 292:501-504. However, these strategies are limited to *recoding* the genetic code rather than *expanding* the genetic code and lead to varying degrees of substitution of one of the common twenty amino acids with an unnatural amino acid.

[0007]   Later it was shown that unnatural amino acids could be site-specifically incorporated into proteins *in vitro* by the addition of chemically aminoacylated orthogonal amber suppressor tRNAs to an *in vitro* transcription/translation reaction. *See, e.g.,* Noren, C. J., et al. (1989) Science 244:182-188; Bain, J. D., et al., (1989) J. Am. Chem. Soc. 111:8013-8014; Dougherty, D. A. (2000) Curr. Opin. Chem. Biol. 4, 645-652; Cornish, V. W., et al. (1995) Angew. Chem., Int. Ed. 34:621-633; J. A. Ellman, et al., (1992), Science 255:197-200; and, D. Mendel, et al., (1995), Annual Review of Biophysics and

Biomolecular Structure 24:435-462. These studies show that the ribosome and translation factors are compatible with a large number of unnatural amino acids, even those with unusual structures. Unfortunately, the chemical aminoacylation of tRNAs is difficult, and the stoichiometric nature of this process severely limited the amount of protein that could be

5   generated.

[0008]   Unnatural amino acids have been microinjected into cells. For example, unnatural amino acids were introduced into the nicotinic acetylcholine receptor in Xenopus oocytes (e.g., M.W. Nowak, et al. (1998), *In vivo incorporation of unnatural amino acids into ion channels in Xenopus oocyte expression system*, Method Enzymol. 293:504-529) by

10   microinjection of a chemically misacylated Tetrahymena thermophila tRNA (e.g., M.E. Saks, et al. (1996), *An engineered Tetrahymena tRNAGln for in vivo incorporation of unnatural amino acids into proteins by nonsense suppression*, J. Biol. Chem. 271:23169-23175), and the relevant mRNA. *See, also*, D.A. Dougherty (2000), *Unnatural amino acids as probes of protein structure and function*, Curr. Opin. Chem. Biol. 4:645-652.

15   Unfortunately, this methodology is limited to proteins in cells that can be microinjected, and, because the relevant tRNA is chemically acylated in vitro, and cannot be re-acylated, the yields of protein are very low.

[0009]   To overcome these limitations, new components, e.g., orthogonal tRNAs, orthogonal aminoacyl-tRNA synthetases and pairs thereof, were added to the protein

20   biosynthetic machinery of the prokaryote *Escherichia coli* (*E. coli*) (see e.g., L. Wang, et al., (2001), Science 292:498-500), which allowed genetic encoding of unnatural amino acids *in vivo*. A number of new amino acids with novel chemical, physical or biological properties, including photoaffinity labels and photoisomerizable amino acids, photocrosslinking amino acids (*see, e.g.,* Chin, J. W., et al. (2002) Proc. Natl. Acad. Sci. U.

25   S. A. 99:11020-11024; and, Chin, J. W., et al., (2002) J. Am. Chem. Soc. 124:9026-9027), keto amino acids (*see, e.g.,* Wang, L., et al., (2003) Proc. Natl. Acad. Sci. U. S. A. 100:56-61), heavy atom containing amino acids, and glycosylated amino acids have been incorporated efficiently and with high fidelity into proteins in *E. coli* in response to, e.g., the amber codon (TAG), using this methodology.

30   [0010]   Several other orthogonal pairs have been reported. Glutaminyl (*see, e.g.,* Liu, D. R., and Schultz, P. G. (1999) Proc. Natl. Acad. Sci. U. S. A. 96:4780-4785), aspartyl (*see, e.g.,* Pastrnak, M., et al., (2000) Helv. Chim. Acta 83:2277-2286), and tyrosyl (*see,*

e.g., Ohno, S., et al., (1998) J. Biochem. (Tokyo, Jpn.) 124:1065-1068; and, Kowal, A. K., et al., (2001) Proc. Natl. Acad. Sci. U. S. A. 98:2268-2273) systems derived from *S. cerevisiae* tRNAs and synthetases have been described for the potential incorporation of unnatural amino acids in *E. coli*. Systems derived from the *E. coli* glutaminyl (*see, e.g.,*

5   Kowal, A. K., et al., (2001) Proc. Natl. Acad. Sci. U. S. A. 98:2268-2273) and tyrosyl (*see, e.g.,* Edwards, H., and Schimmel, P. (1990) Mol. Cell. Biol. 10:1633-1641) synthetase have been described for use in *S. cerevisiae*. The *E. coli* tyrosyl system has been used for the incorporation of 3-iodo-L-tyrosine *in vivo*, in mammalian cells. *See,* Sakamoto, K., et al., (2002) Nucleic Acids Res. 30:4692-4699. Typically, these systems have made use of the

10   amber stop codon. To further expand the genetic code, there is a need to develop improved and/or additional components of the biosynthetic machinery, e.g., additional orthogonal tRNAs, orthogonal aminoacyl-tRNA synthetases, and/or unique codons. This invention fulfills these and other needs, as will be apparent upon review of the following disclosure.

SUMMARY OF THE INVENTION

15   **[0011]**   To expand the genetic code, the invention provides compositions of and methods for producing orthogonal leucyl-tRNAs, orthogonal leucyl aminoacyl-tRNA synthetases and pairs thereof. These translational components can be used to incorporate a selected amino acid in a specific position in a growing polypeptide chain (during nucleic acid translation) in response to a selector codon.

20   **[0012]**   Compositions of the invention include a composition comprising an orthogonal leucyl-tRNA (leucyl-O-tRNA), where the leucyl O-tRNA comprises an anticodon loop comprising a CU(X)$_n$ XXXAA sequence, and comprises at least about a 25% suppression activity in presence of a cognate synthetase in response to a selector codon as compared to a comparable control (e.g., in the absence of the selector codon). In one

25   embodiment, the selector codon is an amber codon, and the leucyl O-tRNA comprises a stem region comprising matched base pairs and a conserved discriminator base at position 73. This position is indicated in Figure 4, Panel A. In one aspect, the CU(X)$_n$ XXXAA sequence comprises CUCUAAA sequence and n=0. In another aspect, the leucyl O-tRNA comprises a C:G base pair at position 3:70.

30   **[0013]**   In one embodiment, the selector codon is a four-base codon and the leucyl O-tRNA comprises a first pair selected from U28:A42, G28:C42 and/or C28:G42, and a

second pair selected from G:49:C65 or C49:G65, where the numbering corresponds to that indicated in Figure 4, Panel A. In one aspect, the CU(X)$_n$XXXAA sequence comprises a CUUCCUAA sequence and n=1. In another aspect, the first pair is C28:G42 and the second pair is C49:G65. In one embodiment, the CU(X)$_n$XXXAA sequence comprises a CUUCAAA sequence and n=0, and the selector codon is an opal codon.

[0014] A composition comprising a leucyl O-tRNA can further include an orthogonal leucyl aminoacyl-tRNA synthetase (leucyl O-RS), where the leucyl O-RS preferentially aminoacylates the leucyl O-tRNA with a selected amino acid. In certain embodiments, a composition including a leucyl O-tRNA can further include a (e.g., in vitro or in vivo) translation system.

[0015] A composition of the invention also includes a cell (e.g., a non-eukaryotic cell (e.g., an *E. coli* cell), or a eukaryotic cell) comprising a translation system. The translation system includes an orthogonal leucyl-tRNA (leucyl-O-tRNA), where the leucyl-O-tRNA comprises at least about a 25% suppression activity in presence of a cognate synthetase in response to a selector codon as compared to a control lacking the selector codon; an orthogonal aminoacyl-leucyl-tRNA synthetase (leucyl-O-RS); and, a first selected amino acid. In these cells, the leucyl O-tRNA comprises an anticodon loop comprising a CU(X)$_n$XXXAA sequence and recognizes the first selector codon and the leucyl O-RS preferentially aminoacylates the leucyl O-tRNA with the first selected amino acid. In some embodiments, the cell translation system comprises a leucyl-O-tRNA and cognate synthetase, or a conservative variant thereof, where these components are at least 50% as effective at suppressing a selector codon as a leucyl O-tRNA of SEQ ID NO: 3, 6, 7 or 12, in combination with a cognate synthetase.

[0016] In certain embodiments, the cell can further include an additional different O-tRNA/O-RS pair and a second selected amino acid, where the O-tRNA recognizes a second selector codon and the O-RS preferentially aminoacylates the O-tRNA with the second selected amino acid. In one embodiment, the cell further comprises a nucleic acid that comprises a polynucleotide that encodes a polypeptide of interest, where the polynucleotide comprises/encodes a selector codon that is recognized by the leucyl O-tRNA.

[0017]   In one embodiment, an *E. coli* cell includes an orthogonal leucyl-tRNA (leucyl-O-tRNA), where the leucyl-O-tRNA comprises at least about a 25% suppression activity in presence of a cognate synthetase in response to a selector codon as compared to a control lacking the selector codon; and an orthogonal leucyl aminoacyl-tRNA synthetase (leucyl-O-RS), where the O-RS preferentially aminoacylates the O-tRNA with a selected amino acid. The E. coli cell also includes the selected amino acid, and, a nucleic acid that comprises a polynucleotide that encodes a polypeptide of interest, where the polynucleotide comprises a selector codon that is recognized by the leucyl O-tRNA. In one example, the leucyl O-tRNA is derived from *Halobacterium sp NRC-1* and the leucyl O-RS is derived from *Methanobacterium thermoaautotropicum.*

[0018]   In certain embodiments of the invention, a leucyl O-tRNA of the invention comprises or is encoded by a polynucleotide sequence as set forth in any one of SEQ ID NO.: 3, 6, 7 or 12, or a complementary polynucleotide sequence thereof. In some embodiments, the leucyl-O-tRNA and cognate synthetase, or a conservative variant thereof, are at least 50% as effective at suppressing a selector codon as a leucyl O-tRNA of SEQ ID NO: 3, 6, 7 or 12, in combination with a cognate synthetase. In the case of tRNA molecules, thymine (t) is, of course, replaced by uracil (u). In certain embodiments, a leucyl O-RS comprises an amino acid sequence as set forth in any one of SEQ ID NO.: 15 or 16, or a conservative variation thereof. In one embodiment, the leucyl O-RS or a portion thereof is encoded by a polynucleotide sequence as set forth in any one of SEQ ID NO.: 13 or 14, a conservative variant of SEQ ID NO: 13 or 14, or a complementary polynucleotide sequence thereof.

[0019]   The leucyl O-tRNA and/or the leucyl O-RS of the invention can be derived from any of a variety of organisms (e.g., both eukaryotic and non-eukaryotic organisms). For example, the leucyl O-tRNA is derived from an archael tRNA (e.g., from *Halobacterium sp NRC-1*) and/or the leucyl O-RS is derived from a non-eukaryotic organism (e.g., *Methanobacterium thermoaautotropicum*).

[0020]   Polynucleotides are also a feature of the invention. A polynucleotide of the invention includes a polynucleotide comprising a nucleotide sequence as set forth in any one of SEQ ID NO.: 1-2, 4-7, 12, and/or is complementary to or that encodes a polynucleotide sequence of the above. A polynucleotide of the invention also includes a nucleic acid that hybridizes to a polynucleotide described above, under highly stringent

conditions over substantially the entire length of the nucleic acid. A polynucleotide of the invention also includes a polynucleotide that is, e.g., at least 75%, at least 80%, at least 90%, at least 95%, at least 98% or more identical to that of a naturally occurring leucyl tRNA or a consensus sequence of multiple naturally occurring leucyl tRNAs, e.g., the laucyl

5   tRNA of SEQ ID NO: 12, and comprises an anticodon loop comprising a $CU(X)_n XXXAA$ sequence, an stem region lacking noncanonical base pairs and a conserved discriminator base at position 73. A polynucleotide of the invention also includes a polynucleotide that is, e.g., at least 75%, at least 80%, at least 90%, at least 95%, at least 98% or more identical to that of a naturally occurring leucyl tRNA and comprises an anticodon loop comprising a

10   CUUCCUAA sequence, a first pair selected from T28:A42, G28:C42 and/or C28:G42, and a second pair selected from G:49:C65 or C49:G65, where the numbering corresponds to that indicated in Figure 4, Panel A. Polynucleotides that are, e.g., at least 80%, at least 90%, at least 95%, at least 98% or more identical to any of the above and/or a polynucleotide comprising a conservative variation of any the above or in Table 3 are also polynucleotides

15   of the invention.

[0021]   Vectors comprising or encoding a polynucleotide of the invention are also a feature of the invention. For example, a vector optionally includes any of: a plasmid, a cosmid, a phage, a virus, an expression vector, and/or the like. A cell comprising a vector of the invention is also a feature of the invention.

20   [0022]   Methods of producing an orthogonal tRNA (O-tRNA), e.g., a leucyl O-tRNA, are also a feature of the invention. An O-tRNA, e.g., a leucyl O-tRNA, produced by the method is also a feature of the invention. For example, a method includes mutating an anticodon loop on members of a pool of tRNAs (e.g., pool of leucyl tRNAs) to allow recognition of a selector codon, thereby providing a plurality of potential O-tRNAs; and

25   analyzing secondary structure of at least one member of the plurality potential O-tRNA to identify non-canonical base pairs in the secondary structure, and, optionally, mutating the non-canonical base pairs (e.g., mutating the non-canonical base pairs to canonical base pairs). In one embodiment, the non-canonical base pairs are located in stem region of the secondary structure. A population of cells of a first species, where the cells individually

30   comprise at least one member of the plurality of potential O-tRNAs are subjected to a negative selection, thereby eliminating cells that comprise a member of the plurality of potential O-tRNAs that is aminoacylated by an aminoacyl-tRNA synthetase (RS) that is

endogenous to the cell, and providing a pool of tRNAs that are orthogonal to the cell of the first species. In certain embodiments, the selector codon includes an amber codon, an opal codon, a four base codon, etc. The method can further include adding an additional sequence (CCA) to a 3' terminus of each of the pool of tRNAs and/or measuring

5   suppression activity.

[0023]   In one embodiment, the pool of tRNAs is obtained by aligning a plurality of tRNA sequences; determining a consensus sequence; and generating a library of mutant tRNAs using the consensus sequence, where the pool of tRNAs comprise the library of mutant tRNAs.

10   [0024]   In certain embodiments, the subjecting step comprises a polynucleotide that encodes a negative selection marker. In one embodiment, the polynucleotide that encodes the negative selection marker comprises at least one selector codon. For example, a negative selection marker includes, but is not limited to, β-lactamase, β-galactosidase, and/or the like. In certain embodiments, the negative selection marker fluoresces or

15   catalyzes a luminescent reaction in the presence of a suitable reactant. In another embodiment, a product of the negative selection marker is detected by fluorescence-activated cell sorting (FACS) or by luminescence. Optionally, the negative selection marker includes an affinity based screening marker. In certain embodiments, the subjecting step comprises growing the population of cells in the presence of a selective agent (e.g., an

20   antibiotic, such as ampicillin).

[0025]   In certain embodiments, the method further comprises subjecting to positive selection a second population of cells of the first species. The cells comprise a member of the pool of tRNAs that are orthogonal to the cell of the first species, a cognate aminoacyl-tRNA synthetase, and a positive selection marker. Cells are selected/screened for cells that

25   comprise a member of the pool of tRNAs that is aminoacylated by the cognate aminoacyl-tRNA synthetase and that shows a desired response in the presence of the positive selection marker, thereby providing an O-tRNA.

[0026]   Methods for identifying an orthogonal aminoacyl-tRNA synthetase (O-RS), e.g., a leucyl O-RS, for use with an O-tRNA, e.g., a leucyl O-tRNA, are also a feature of the

30   invention. For example, a method includes subjecting to positive selection a population of cells of a first species, where the cells each comprise: 1) a member of a plurality of

aminoacyl-tRNA synthetases (RSs), where the plurality of RSs comprise mutant RSs, RSs derived from a species other than the first species or both mutant RSs and RSs derived from a species other than the first species; 2) the orthogonal tRNA (O-tRNA) (e.g., from a species other than the first species, from at least a second species, etc.); and 3) a

5   polynucleotide that encodes a positive selection marker and comprises at least one selector codon. In one embodiment, the plurality of RSs comprises leucyl RSs. In certain embodiments, the O-tRNA comprises a leucyl O-tRNA (e.g., where leucyl O-tRNA includes at least about a 25% suppression activity in presence of a cognate synthetase in response to a selector codon as compared to a control lacking the cognate synthetase).

10   [0027]   Cells are selected or screened for those that show an enhancement in suppression efficiency compared to cells lacking or having a reduced amount of the member of the plurality of RSs. These selected/screened cells comprise an active RS that aminoacylates the O-tRNA. The level of aminoacylation (in vitro or in vivo) by the active RS of a first set of tRNAs from the first species is compared to the level of aminoacylation

15   (in vitro or in vivo) by the active RS of a second set of tRNAs from a second species; where the level of aminoacylation is determined by a detectable substance (e.g., a labeled amino acid). The active RS that more efficiently aminoacylates the second set of tRNAs compared to the first set of tRNAs is selected, thereby providing the orthogonal aminoacyl-tRNA synthetase, e.g., leucyl O-RS, for use with the O-tRNA, e.g., the leucyl O-tRNA. An

20   orthogonal aminoacyl-tRNA synthetase identified by the method is also a feature of the invention.

[0028]   Methods of producing a protein in a cell with a selected amino acid at a specified position are also a feature of the invention. For example, a method includes growing, in an appropriate medium, a cell, where the cell comprises a nucleic acid that

25   comprises at least one selector codon and encodes a protein; and, providing the selected amino acid. The cell further comprises: an orthogonal leucyl-tRNA (leucyl-O-tRNA) that functions in the cell and recognizes the selector codon; and, an orthogonal leucyl aminoacyl-tRNA synthetase (leucyl O-RS) that preferentially aminoacylates the leucyl-O-tRNA with the selected amino acid. Typically, the leucyl-O-tRNA comprises at least about

30   a 25% suppression activity in presence of a cognate synthetase in response to a selector codon as compared to a control lacking the cognate synthetase. A protein produced by this method is also a feature of the invention.

DEFINITIONS

[0029]   Before describing the invention in detail, it is to be understood that this
invention is not limited to particular biological systems, which can, of course, vary. It is
also to be understood that the terminology used herein is for the purpose of describing
particular embodiments only, and is not intended to be limiting. As used in this
specification and the appended claims, the singular forms "a", "an" and "the" include plural
referents unless the content clearly dictates otherwise. Thus, for example, reference to "a
cell" includes a combination of two or more cells; reference to "bacteria" includes mixtures
of bacteria, and the like.

[0030]   Unless defined herein and below in the reminder of the specification, all
technical and scientific terms used herein have the same meaning as commonly understood
by one of ordinary skill in the art to which the invention pertains.

[0031]   Orthogonal leucyl-tRNA: As used herein, an orthogonal leucyl-tRNA
(leucyl-O-tRNA) is a tRNA that is orthogonal to a translation system of interest, where the
tRNA is: (1) identical or substantially similar to a naturally occurring leucyl tRNA, (2)
derived from a naturally occurring leucyl tRNA by natural or artificial mutagenesis (3)
derived by any process that takes a sequence of a wild-type or mutant leucyl tRNA
sequence of (1) or (2) into account, (4) homologous to a wild-type or mutant leucyl tRNA;
(5) homologous to any example tRNA that is designated as a substrate for a leucyl tRNA
synthetase in Table 3, or (6) a conservative variant of any example tRNA that is designated
as a substrate for a leucyl tRNA synthetase in Table 3. The leucyl tRNA can exist charged
with a amino acid, or in an uncharged state. It is also to be understood that a "leucyl-O-
tRNA" optionally is charged (aminoacylated) by a cognate synthetase with an amino acid
other than leucine. Indeed, it will be appreciated that a leucyl-O-tRNA of the invention is
advantageously used to insert essentially any amino acid, whether natural or artificial, into a
growing polypeptide, during translation, in response to a selector codon.

[0032]   Orthogonal leucyl amino acid synthetase: As used herein, an orthogonal
leucyl amino acid synthetase (leucyl O-RS) is an enzyme that preferentially aminoacylates
the leucyl-O-tRNA with an amino acid in a translation system of interest. The amino acid
that the leucyl O-RS loads onto the leucyl O-tRNA can be any amino acid, whether natural
or artificial, and is not limited herein. The synthetase is optionally the same as or
homologous to a naturally occurring leucyl amino acid synthetase, or the same as or

homologous to a synthetase designated as a leucyl O-RS in Table 3. For example, the leucyl O-RS can be a conservative variant of a leucyl O-RS of Table 3, and/or can be at least 50%, 60%, 70%, 80%, 90%, 95%, 98%, 99% or more identical in sequence to a leucyl O-RS of Table 3.

5   [0033]   Homologous: Proteins and/or protein sequences are "homologous" when they are derived, naturally or artificially, from a common ancestral protein or protein sequence. Similarly, nucleic acids and/or nucleic acid sequences are homologous when they are derived, naturally or artificially, from a common ancestral nucleic acid or nucleic acid sequence. For example, any naturally occurring nucleic acid can be modified by any

10   available mutagenesis method to include one or more selector codon. When expressed, this mutagenized nucleic acid encodes a polypeptide comprising one or more selected amino acid, e.g. unnatural amino acid. The mutation process can, of course, additionally alter one or more standard codon, thereby changing one or more standard amino acid in the resulting mutant protein as well. Homology is generally inferred from sequence similarity between

15   two or more nucleic acids or proteins (or sequences thereof). The precise percentage of similarity between sequences that is useful in establishing homology varies with the nucleic acid and protein at issue, but as little as 25% sequence similarity is routinely used to establish homology. Higher levels of sequence similarity, e.g., 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, or 99% or more, can also be used to establish homology. Methods for

20   determining sequence similarity percentages (e.g., BLASTP and BLASTN using default parameters) are described herein and are generally available.

       [0034]   Orthogonal: As used herein, the term "orthogonal" refers to a molecule (e.g., an orthogonal tRNA (O-tRNA) and/or an orthogonal aminoacyl tRNA synthetase (O-RS)) that functions with endogenous components of a cell with reduced efficiency as compared

25   to a corresponding molecule that is endogenous to the cell or translation system, or that fails to function with endogenous components of the cell. In the context of tRNAs and aminoacyl-tRNA synthetases, orthogonal refers to an inability or reduced efficiency, e.g., less than 20 % efficiency, less than 10 % efficiency, less than 5 % efficiency, or less than 1% efficiency, of an orthogonal tRNA to function with an endogenous tRNA synthetase

30   compared to the ability of an endogenous tRNA to function with the endogenous tRNA synthetase; or of an orthogonal aminoacyl-tRNA synthetase to function with an endogenous tRNA compared to the ability of an endogenous tRNA synthetase to function with the

endogenous tRNA. The orthogonal molecule lacks a functionally normal endogenous complementary molecule in the cell. For example, an orthogonal tRNA in a cell is aminoacylated by any endogenous RS of the cell with reduced or even undetectable efficiency, when compared to aminoacylation of an endogenous tRNA by the endogenous

5   RS. In another example, an orthogonal RS aminoacylates any endogenous tRNA in a cell of interest with reduced or even undetectable efficiency, as compared to aminoacylation of the endogenous tRNA by an endogenous RS. A second orthogonal molecule can be introduced into the cell that functions with the first orthogonal molecule. For example, an orthogonal tRNA/RS pair includes introduced complementary components that function together in the

10   cell with an efficiency (e.g., 45 % efficiency, 50% efficiency, 60% efficiency, 70% efficiency, 75% efficiency, 80% efficiency, 90% efficiency, 95% efficiency, or 99% or more efficiency) as compared to that of a control, e.g., a corresponding tRNA/RS endogenous pair, or an active orthogonal pair (e.g., a tyrosyl orthogonal tRNA/RS pair).

[0035]   Cognate: The term "cognate" refers to components that function together,

15   e.g., a leucyl tRNA and a leucyl aminoacyl-tRNA synthetase. The components can also be referred to as being complementary.

[0036]   Preferentially aminoacylates: The term "preferentially aminoacylates" refers to an efficiency, e.g., 70 % efficient, 75 % efficient, 85% efficient, 90% efficient, 95 % efficient, or 99% or more efficient, at which an O-RS aminoacylates an O-tRNA with a

20   selected amino acid, e.g., an unnatural amino acid, as compared to the O-RS aminoacylating a naturally occurring tRNA or a starting material used to generate the O-tRNA.

[0037]   Selector codon: The term "selector codon" refers to codons recognized by the O-tRNA in the translation process and not recognized by an endogenous tRNA. The O-tRNA anticodon loop recognizes the selector codon on the mRNA and incorporates its

25   amino acid, e.g., a selected amino acid, such as an unnatural amino acid, at this site in the polypeptide. Selector codons can include, e.g., nonsense codons, such as, stop codons, e.g., amber, ochre, and opal codons; four or more base codons; rare codons; codons derived from natural or unnatural base pairs and/or the like.

[0038]   Suppressor tRNA: A suppressor tRNA is a tRNA that alters the reading of a

30   messenger RNA (mRNA) in a given translation system, e.g., by providing a mechanism for incorporating an amino acid into a polypeptide chain in response to a selector codon. For

example, a suppressor tRNA can read through, e.g., a stop codon, a four base codon, or a rare codon.

[0039]   Suppression activity: As used herein, the term "suppression activity" refers, in general, to the ability of a tRNA (e.g., a suppressor tRNA) to allow translational read-through of a codon (e.g. a selector codon that is an amber codon or a 4-or-more base codon) that would otherwise result in the termination of translation or mistranslation (e.g., frame-shifting). Suppression activity of a suppressor tRNA can be expressed as a percentage of translational read-through observed compared to a second suppressor tRNA, or as compared to a control system, e.g., a control system lacking an O-RS.

[0040]   The present invention provides various means by which suppression activity can be quantitated. Percent suppression of a particular OtRNA and ORS against a selector codon (e.g., an amber codon) of interest refers to the percentage of activity of a given expressed test marker (e.g., LacZ), that includes a selector codon, in a nucleic acid encoding the expressed test marker, in a translation system of interest, where the translation system of interest includes an O-RS and an O-tRNA, as compared to a positive control construct, where the positive control lacks the O-tRNA, the O-RS and the selector codon. Thus, for example, if an active positive control marker construct that lacks a selector codon has an observed activity of X in a given translation system, in units relevant to the marker assay at issue, then percent suppression of a test construct comprising the selector codon is the percentage of X that the test marker construct displays under essentially the same environmental conditions as the positive control marker was expressed under, except that the test marker construct is expressed in a translation system that also includes the O-tRNA and the O-RS. Typically, the translation system expressing the test marker also includes an amino acid that is recognized by the O-RS and O-tRNA. Optionally, the percent suppression measurement can be refined by comparison of the test marker to a "background" or "negative" control marker construct, which includes the same selector codon as the test marker, but in a system that does not include the O-tRNA, O-RS and/or relevant amino acid recognized by the O-tRNA and/or O-RS. This negative control is useful in normalizing percent suppression measurements to account for background signal effects from the marker in the translation system of interest.

[0041]   Suppression efficiency can be determined by any of a number of assays known in the art. For example, a β-galactosidase reporter assay can be used, e.g., a

derivatized lacZ plasmid (where the construct has a selector codon in the lacZ nucleic acid sequence) is introduced into cells from an appropriate organism (e.g., an organism where the orthogonal components can be used) along with plasmid comprising an O-tRNA of the invention. A cognate synthetase can also be introduced (either as a polypeptide or a

5   polynucleotide that encodes the cognate synthetase when expressed). The cells are grown in media to a desired density, e.g., to an OD_{600} of about 0.5, and β-galactosidase assays are performed, e.g., using the BetaFluor™ β-Galactosidase Assay Kit (Novagen). Percent suppression can be calculated as the percentage of activity for a sample relative to a comparable control, e.g., the value observed from the derivatived lacZ construct, where the

10   construct has a corresponding sense codon at desired position rather than a selector codon.

[0042]   Translation system: The term "translation system" refers to the components that incorporate an amino acid into a growing polypeptide chain (protein). Components of a translation system can include, e.g., ribosomes, tRNAs, synthetases, mRNA and the like. The O-tRNA and/or O-RS of the invention can be added to or be a part of an *in vitro* or *in*

15   *vivo* translation system, e.g., in a non-eukaryotic cell, e.g., a bacterium (such as *E coli*), or in a eukaryotic cell, e.g., a yeast cell, a mammalian cell, a plant cell, an algae cell, a fungus cell, an insect cell, and/or the like.

[0043]   Selected amino acid: The term "selected amino acid" refers to any desired naturally occurring amino acid or unnatural amino acid. As used herein, the term

20   "unnatural amino acid" refers to any amino acid, modified amino acid, and/or amino acid analogue that is not one of the 20 common naturally occurring amino acids or seleno cysteine or pyrolysine.

[0044]   Derived from: As used herein, the term "derived from" refers to a component that is isolated from or made using a specified molecule or organism, or information from

25   the specified molecule or organism.

[0045]   Positive selection or screening marker: As used herein, the term "positive selection or screening marker" refers to a marker that, when present, e.g., expressed, activated, or the like, results in identification of a cell with the positive selection marker from those without the positive selection marker.

30   [0046]   Negative selection or screening marker: As used herein, the term "negative selection or screening marker" refers to a marker that, when present, e.g., expressed,

activated or the like, allows identification of a cell that does not possess a specified property (e.g., as compared to a cell that does possess the property).

[0047]   Reporter: As used herein, the term "reporter" refers to a component that can be used to identify and/or select target components of a system of interest. For example, a reporter can include a protein, e.g., an enzyme, that confers antibiotic resistance or sensitivity (e.g., β-lactamase, chloramphenicol acetyltransferase (CAT), and the like), a fluorescent screening marker (e.g., green fluorescent protein (e.g., (GFP), YFP, EGFP, RFP), a luminescent marker (e.g., a firefly luciferase protein), an affinity based screening marker, or positive or negative selectable marker genes such as lacZ, β-gal/lacZ (β-galactosidase), Adh (alcohol dehydrogenase), his3, ura3, leu2, lys2, or the like.

[0048]   Eukaryote: As used herein, the term "eukaryote" refers to organisms belonging to the phylogenetic domain Eucarya, such as animals (e.g., mammals, insects, reptiles, birds, etc.), ciliates, plants (e.g., monocots, dicots, algae, etc.), fungi, yeasts, flagellates, microsporidia, protists, etc.

[0049]   Non-eukaryote: As used herein, the term "non-eukaryote" refers to non-eukaryotic organisms. For example, a non-eukaryotic organism can belong to the Eubacteria (e.g., *Escherichia coli, Thermus thermophilus, Bacillus stearothermophilus*, etc.) phylogenetic domain, or the Archaea (e.g., *Methanococcus jannaschii, Methanobacterium thermoautotrophicum, Halobacterium* such as *Haloferax volcanii* and *Halobacterium* species *NRC-1, Archaeoglobus fulgidus, Pyrococcus furiosus, Pyrococcus horikoshii, Aeuropyrum pernix,* etc.) phylogenetic domains.

[0050]   Conservative variant: The term "conservative variant" in reference to a translation component such as an O-tRNA or O-RS refers to a translation component that has a substantailly similar activity as the component on which the conservative variant is similar to, e.g., an O-tRNA or O-RS, but has variations in the sequence as compared to the base component. For example, an O-RS will aminoacylate a complementary O-tRNA or a conservative variant O-tRNA with a selected amino acid, e.g., an unnatural amino acid, although the O-tRNA and the conservative variant O-tRNA do not have the same sequence. The conservative variant can have, e.g., one variation, two variations, three variations, four variations, or five or more variations in its sequence, as long as the conservative variant

functionally interacts with a corresponding O-tRNA or O-RS in substantailly the same manner as the non-variant form. ·

[0051]   Selection or screening agent: As used herein, the term "selection or screening agent" refers to an agent that, when present, allows for selection/screening of certain components from a population. For example, a selection or screening agent can be, but is not limited to, e.g., a nutrient, an antibiotic, a wavelength of light, an antibody, an expressed polynucleotide, or the like. The selection agent can be varied, e.g., by concentration, intensity, etc.

[0052]   Encode: As used herein, the term "encode" refers to any process whereby the information in a polymeric macromolecule or sequence string is used to direct the production of a second molecule or sequence string that is different from the first molecule or sequence string. As used herein, the term is used broadly, and can have a variety of applications. In one aspect, the term "encode" describes the process of semi-conservative DNA replication, where one strand of a double-stranded DNA molecule is used as a template to encode a newly synthesized complementary sister strand by a DNA-dependent DNA polymerase.

[0053]   In another aspect, the term "encode" refers to any process whereby the information in one molecule is used to direct the production of a second molecule that has a different chemical nature from the first molecule. For example, a DNA molecule can encode an RNA molecule (e.g., by the process of transcription incorporating a DNA-dependent RNA polymerase enzyme). Also, an RNA molecule can encode a polypeptide, as in the process of translation. When used to describe the process of translation, the term "encode" also extends to the triplet codon that encodes an amino acid. In some aspects, an RNA molecule can encode a DNA molecule, e.g., by the process of reverse transcription incorporating an RNA-dependent DNA polymerase. In another aspect, a DNA molecule can encode a polypeptide, where it is understood that "encode" as used in that case incorporates both the processes of transcription and translation.

BRIEF DESCRIPTION OF THE FIGURES

[0054]   Figure 1, Panels A, B and C schematically illustrate leucyl tRNAs and synthetases, and their phylogenetic relationships. Panel A provides a ClustalW analysis of aminoacyl-tRNA synthetases, where Archaeal tRNA synthetases are labeled using a dashed

line, prokaryotic using a solid line, and eukaryotic sequences using a dotted line. This analysis reveals the halobacterial synthetase to be unusual in its homology to prokaryotic rather than archaeal and eukaryotic synthetases. **Panel B** provides a ClustalW analysis of Halobacterial tRNAs which all share high homology to other archaeal tRNAs.

5   Dendrograms were generated using the program PhyloDraw. **Panel C** provides a sequence alignment of multiple sequences of the family of archaeal leucyl tRNAs examined as potential orthogonal suppressors. Sequences examined as potential amber suppressors by changing the anticodon (boxed) to CUA are shown in bold as is the consensus sequence. The highly conserved positions G37 and A73 are indicated with underlining.

10 **[0055]**   **Figure 2** provides a histogram showing the identification of a leucyl orthogonal pair. The suppression efficiency of seven synthetases expressed with 5 orthogonal amber suppressor reporter constructs was measured using a β-lactamase amber suppression assay.

[0056]   **Figure 3, Panels A** and **B** provide graphs illustrating aminoacylation *in vitro*

15 by archaeal leucyl-tRNA synthetases. **Panel A** illustrates charging of crude total halobacterial tRNA determined by aminoacylation assays with [^{3}H] leucine by AfLRS ($\blacksquare$), MjLRS ($\bullet$), MtLRS ($\blacktriangle$), EcLRS ($\blacklozenge$), and no synthetase ($\square$). **Panel B** illustrates charging of crude total *E. coli* tRNA.

[0057]   **Figure 4, Panels A** and **B** illustrates the optimization of suppressor tRNAs.

20 **Panel A** illustrates regions (shown in boxes) of the halobacterial orthogonal tRNA subjected to mutagenesis in an effort to improve the efficiency or selectivity of TAG and AGGA suppressor tRNAs. **Panel B** illustrates that active mutant TAG suppressors identified by positive selection conserve A73. Less cross-reactive mutants identified by a double-sieve selection strategy conserve a C3:G70 base pair. The most active and selective

25 suppressor tRNA is shown with double boxes.

[0058]   **Figure 5** illustrates a consensus-derived frameshift suppressor. A consensus sequence was obtained by multiple sequence alignment of all known archaeal leucyl tRNAs, and the anticodon loop is changed to UCUCCUAA. The variations observed for tRNAs identified by selection are shown in boxes. The most active mutations are shown with

30 double boxes.

DETAILED DESCRIPTION

[0059]   In order to add additional unnatural amino acids to the genetic code *in vivo*, "orthogonal pairs" of an aminoacyl-tRNA synthetase and a tRNA are needed that can function efficiently in the translational machinery. Desired characteristics of the orthogonal pairs include tRNA that decode or recognize only a specific new codon, e.g., a selector codon, that is not decoded by any endogenous tRNA, and aminoacyl-tRNA synthetases that preferentially aminoacylate (or charge) its cognate tRNA with only a specific selected amino acid, e.g., an unnatural amino acid. The O-tRNA is also not typically aminoacylated by endogenous synthetases. For example, in *E. coli*, an orthogonal pair will include an aminoacyl-tRNA synthetase that does not significantly cross-react with any of the endogenous tRNA, which there are 40 in *E. coli*, and an orthogonal tRNA that is not significantly aminoacylated by any of the endogenous synthetases, e.g., of which there are 21 in *E. coli*.

[0060]   The O-tRNA is capable of mediating incorporation of a selected amino acid into a protein that is encoded by a polynucleotide, which comprises a selector codon that is recognized by the O-tRNA, e.g., in vivo. The anticodon loop of the O-tRNA recognizes the selector codon on an mRNA and incorporates its amino acid, e.g., a selected amino acid, such as an unnatural amino acid, at this site in the polypeptide. Any of a number of selector codons can be used with the invention. For example, selector codons can include, e.g., nonsense codons, such as, stop codons, e.g., amber, ochre, and opal codons; four or more base codons; rare codons; codons derived from natural or unnatural base pairs and/or the like. *See also* the section herein entitled "Selector codon."

[0061]   By using different selector codons, multiple orthogonal tRNA/synthetase pairs can be developed that allow the simultaneous incorporation of multiple selected amino acids, e.g., unnatural amino acids, using these different selector codons. This invention provides compositions of and methods for identifying and producing additional orthogonal tRNA-aminoacyl-tRNA synthetase pairs, e.g., leucyl O-tRNA/leucyl O-RSs, using any of a number of selector codons, e.g., an amber codon, an opal codon, an extended codon (such as a four-base codon), and the like.

ORTHOGONAL LEUCYL tRNA/ ORTHOGONAL LEUCYL AMINOACYL-tRNA
SYNTHETASES AND PAIRS THEREOF

[0062]   Such translation systems of the invention generally comprise cells that
include an orthogonal leucyl tRNA (leucyl O-tRNA), an orthogonal leucyl aminoacyl tRNA
synthetase (leucyl O-RS), and a selected amino acid, e.g., an unnatural amino acid, where
the leucyl O-RS aminoacylates the leucyl O-tRNA with the selected amino acid. An
orthogonal pair of the invention is composed of a leucyl O-tRNA, e.g., a suppressor tRNA,
a frameshift tRNA, or the like, and an leucyl O-RS. The leucyl-O-tRNA recognize a first
selector codon and has at least about a 25% suppression activity in presence of a cognate
synthetase in response to a selector codon as compared to a control lacking the cognate
synthetase. The leucyl O-tRNA also comprises an anticodon loop comprising a $CU(X)_n$
XXXAA sequence. The cell uses the components to incorporate the selected amino acid
into a growing polypeptide chain. For example, a nucleic acid that comprises a
polynucleotide that encodes a polypeptide of interest can also be present, where the
polynucleotide comprises a selector codon that is recognized by the leucyl O-tRNA.The
translation system can also be an in vitro system.

[0063]   Translation systems that are suitable for making proteins that include one or
more selected amino acids, e.g., an unnatural amino acid, are described in International
patent applications WO 2002/086075, entitled "METHODS AND COMPOSITION FOR
THE PRODUCTION OF ORTHOGANOL tRNA-AMINOACYLtRNA SYNTHETASE
PAIRS" and WO 2002/085923, entitled "IN VIVO INCORPORATION OF UNNATURAL
AMINO ACIDS." In addition, see International Application Number PCT/US2004/011786,
filed April 16, 2004. Each of these applications is incorporated herein by reference in its
entirety. These translation systems can be adapted to the present invention by substituting
the leucyl-O-RS and leucyl-O-tRNA provided herein.

[0064]   In certain embodiments, a cell of teh invention, e.g., an *E. coli* cell, includes
such a translation system of the invention. For example, an *E. coli* cell of the invention can
include an orthogonal leucyl-tRNA (leucyl-O-tRNA), where the leucyl-O-tRNA comprises
at least about a 25% suppression activity in presence of a cognate synthetase in response to
a selector codon as compared to a control lacking the cognate synthetase; an orthogonal
leucyl aminoacyl-tRNA synthetase (leucyl-O-RS); a selected amino acid; and, a nucleic

acid that comprises a polynucleotide that encodes a polypeptide of interest, where the polynucleotide comprises a selector codon that is recognized by the leucyl O-tRNA.

[0065]   The invention also features multiple O-tRNA/O-RS pairs in a cell, which allows incorporation of more than one selected amino acid. In certain embodiments, the cell can further include an additional different O-tRNA/O-RS pair and a second selected amino acid, where the O-tRNA recognizes a second selector codon and the O-RS preferentially aminoacylates the O-tRNA with the second selected amino acid. For example, a cell can further comprise, e.g., an amber suppressor tRNA-aminoacyl tRNA synthetase pair derived from the tyrosyl-tRNA synthetase of *Methanococcus jannaschii*.

[0066]   The leucyl O-tRNA and/or the leucyl O-RS can be naturally occurring or can be derived by mutation of a naturally occurring tRNA and/or RS, e.g., which generates libraries of tRNAs and/or libraries of RSs, from a variety of organisms. For example, one strategy of producing an orthogonal leucyl tRNA/leucyl aminoacyl-tRNA synthetase pair involves importing a heterologous tRNA/synthetase pair from, e.g., a source other than the host cell, or multiple sources, into the host cell. The properties of the heterologous synthetase candidate include, e.g., that it does not charge any host cell tRNA, and the properties of the heterologous tRNA candidate include, e.g., that it is not aminoacylated by any host cell synthetase. In addition, the heterologous tRNA is orthogonal to all host cell synthetases.

[0067]   A second strategy for generating an orthogonal pair involves generating mutant libraries from which to screen and/or select a leucyl O-tRNA or leucyl O-RS. These strategies can also be combined.

[0068]   In various embodiments, the leucyl O-tRNA and leucyl O-RS are derived from at least one organism. In another embodiment, the leucyl O-tRNA is derived from a naturally occurring or mutated naturally occurring tRNA from a first organism and the leucyl O-RS is derived from naturally occurring or mutated naturally occurring RS from a second organism. In one embodiment, the first and second organism is different. For example, an orthogonal pair of the invention includes a leucyl-tRNA synthetase derived from *Methanobacterium thermoautotrophicum*, and a leucyl tRNA derived from an archael tRNA (e.g., from *Halobacterium sp. NRC-1*). Alternatively, the first and second organism

are the same. See the section entitled "Sources and Hosts" herein for additional
information.

[0069]   In certain embodiments of the invention, a leucyl O-tRNA of the invention
comprises or is encoded or transcribed by or from a polynucleotide sequence as set forth in
5   any one of SEQ ID NO.: 3, 6, 7 or 12, or a complementary polynucleotide sequence thereof.
In certain embodiments, a leucyl O-RS comprises an amino acid sequence as set forth in
any one of SEQ ID NO.: 15 or 16, or a conservative variation thereof. The leucyl O-RS, or
a portion thereof, can also be encoded or transcribed by or from a polynucleotide sequence
as set forth in any one of SEQ ID NO.: 13 or 14, or a complementary polynucleotide
10   sequence thereof. *See also,* the section entitled "Nucleic Acid and Polypeptide Sequence
and Variants," herein.

Orthogonal tRNA (O-tRNA)

[0070]   An orthogonal leucyl tRNA (leucyl O-tRNA) mediates incorporation of a
selected amino acid into a protein that is encoded by a polynucleotide that comprises a
15   selector codon that is recognized by the leucyl O-tRNA, e.g., in vivo. A leucyl O-tRNA of
the invention comprises an anticodon loop comprising a $CU(X)_n XXXAA$ sequence.

[0071]   The $CU(X)_n XXXAA$ sequence is found in the anticodon loop, where X
refers to any nucleotide, and $(X)_n$ is optionally present. The n refers to a number of bases
the anticodon loop is extended, based on the desired selector codon, e.g., a stop codon
20   (n=0), an extended codon, such as a four- (n=1), five- (n=2), six- (n=3) base pair, etc.

[0072]   In one aspect of the invention, the $CU(X)_n XXXAA$ sequence comprises
CUCUAAA sequence (n=0), typically when the selector codon is an amber codon. In
addition, the leucyl O-tRNA can include a stem region comprising matched base pairs and a
conserved discriminator base (position 73). *See, e.g.,* **Figure 4, Panel B**. This position is
25   indicated in e.g., **Figure 4, Panel A**. The leucyl O-tRNA also optionally includes a C:G
base pair at position 3:70.

[0073]   In one example, the $CU(X)_n XXXAA$ sequence comprises a CUUCCUAA
sequence, typically when the selector codon is a four-base codon. *See, e.g.,* **Figure 5**. The
leucyl O-tRNA can also include a first pair selected from T28:A42, G28:C42 and/or
30   C28:G42, and a second pair selected from G:49:C65 or C49:G65, where the numbering
corresponds to that indicated in **Figure 4, Panel A**. In one embodiment, C28:G42 is the

first pair and C49:G65 is the second pair. When the selector codon is an opal codon, the CU(X)ₙ XXXAA sequence can comprises a CUUCAAA sequence.

[0074]   A leucyl O-tRNA of the invention comprises at least about a 25% suppression activity in presence of a cognate synthetase in response to a selector codon, as

5   compared to a control lacking the cognate synthetase. Suppression activity can be determined by any of a number of assays known in the art. For example, a *β*-galactosidase reporter assay can be used. A derivative of a plasmid that expresses *lacZ* gene under the control of promoter is used, e.g., where the Leu-25 of the peptide VVLQRRDWEN of *lacZ* is replaced by a selector codon, e.g., TAG, TGA, AGGA, etc. codons, or sense codons (as a

10   control) for tyrosine, serine, leucine, etc. The derivatived lacZ plasmid is introduced into cells from an appropriate organism (e.g., an organism where the orthogonal components can be used) along with plasmid comprising a O-tRNA of the invention. A cognate synthetase can also be introduced (either as a polypeptide or a polynucleotide that encodes the cognate synthetase when expressed). The cells are grown in media to a desired density, e.g., to an

15   OD_{600} of about 0.5., and β-galactosidase assays are performed, e.g., using the BetaFluor™ β-Galactosidase Assay Kit (Novagen). Percent suppression is calculated as the percentage of activity for a sample relative to a comparable control, e.g., the value observed from the derivatived lacZ construct, where the construct has a corresponding sense codon at desired position rather than a selector codon.

20   [0075]   Examples of leucyl O-tRNAs of the invention are transcribed from any one of SEQ ID NO.: 1-7 and/or 12. *See*, Table 3 and Example 2, herein, for sequences of exemplary O-RS and O-tRNA molecules. In the tRNA molecule, Thymine (T) is replace with Uracil (U); the tRNAs have the same sequence, except for the usual substitution of U's for T's. One of skill will appreciate that the RNA and DNA versions of a tRNA are often

25   referred to simply by reference to the DNA sequence that corersponds to the RNA form of the tRNA. Any time a DNA form of a tRNA is given, one of skill will easily be able to derive the RNA (or vice versa) by strandard transcription (or reverse transcription). In addition, additional modifications to the bases can be present. The invention also includes conservative variations of leucyl O-tRNA. For example, conservative variations of leucyl

30   O-tRNA include those molecules that function like the leucyl O-tRNA of any one of SEQ ID NO.: 1-7 and 12 and maintain the tRNA L-shaped structure, but do not have the same

sequence (and are other than wild type leucyl tRNA molecules). *See also,* the section herein entitled "Nucleic acids and Polypeptides Sequence and Variants."

[0076]   The composition comprising a leucyl O-tRNA can further include an orthogonal leucyl aminoacyl-tRNA synthetase (leucyl O-RS), where the leucyl O-RS

5   preferentially aminoacylates the leucyl O-tRNA with a selected amino acid (e.g., an unnatural amino acid). In certain embodiments, a composition that includes a leucyl O-tRNA can further include a translation system (e.g., in vitro or in vivo). A nucleic acid that comprises a polynucleotide that encodes a polypeptide of interest, where the polynucleotide comprises a selector codon that is recognized by the leucyl O-tRNA, or a combination of

10   one or more of these can also be present in the cell. *See also,* the section herein entitled "Orthogonal aminoacyl-tRNA synthetases."

[0077]   Methods of producing an orthogonal tRNA (O-tRNA), e.g., a leucyl O-tRNA, are also a feature of the invention. An O-tRNA, e.g., a leucyl O-tRNA, produced by the method is also a feature of the invention. For example, a method includes mutating an

15   anticodon loop of members of a pool of tRNAs (e.g., a pool of leucyl tRNAs) to allow recognition of a selector codon (e.g., an amber codon, an opal codon, a four base codon, etc.), thereby providing a plurality of potential O-tRNAs; and analyzing secondary structure of a member of the plurality potential O-tRNA to identify non-canonical base pairs in the secondary structure, and optionally mutating the non-canonical base pairs (e.g., the non-

20   canonical base pairs are mutated to canonical base pairs). The non-canonical base pairs can be located in stem region of the secondary structure. Typically, a leucyl O-tRNA possesses an improvement of orthogonality for a desired organism compared to the starting material, e.g., the plurality of tRNA sequences, while preserving its affinity towards a desired RS.

[0078]   The methods optionally include analyzing the homology of sequences of

25   tRNAs and/or aminoacyl-tRNA synthetases to determine potential candidates for an O-tRNA, O-RS and/or pairs thereof, that appear to be orthogonal for a specific organism. Computer programs known in the art and described herein can be used for the analysis. In one example, to choose potential orthogonal translational components for use in *E. coli,* a prokaryotic organism, a synthetase and/or a tRNA is chosen that does not display unusual

30   homology to prokaryotic organisms.

[0079]   The pool of tRNAs can also be produced by a consensus strategy. For example, the pool of tRNAs is produced by aligning a plurality of tRNA sequences (see e.g., **Figure 1, Panel C**); determining a consensus sequence (see e.g., **Figure 1, Panel C**); and generating a library of tRNAs using at least a portion, most of, or the entire consensus sequence. For example, a consensus sequence can be compiled with a computer program, e.g., the GCG program *pileup*. Optionally, degenerate positions determined by the program are changed to the most frequent base at those positions. A library is synthesized by techniques known in the art using the consensus sequence. For example, overlap extension of oligonucleotides in which each site of the tRNA gene can be synthesized as a doped mixture of 90% the consensus sequence and 10% a mixture of the other 3 bases can be used to provide the library based on the consensus sequence. Other mixtures can also be used, e.g., 75% the consensus sequence and 25% a mixture of the other 3 bases, 80% the consensus sequence and 20% a mixture of the other 3 bases, 95% the consensus sequence and 5% a mixture of the other 3 bases, etc.

[0080]   The library of mutant tRNAs can be generated using various mutagenesis techniques known in the art. For example, the mutant tRNAs can be generated by site-specific mutations, random point mutations, homologous recombination, DNA shuffling or other recursive mutagenesis methods, chimeric construction or any combination thereof.

[0081]   Additional mutations can be introduced at a specific position(s), e.g., at a nonconservative position(s), or at a conservative position, at a randomized position(s), or a combination of both in a desired loop or region of a tRNA, e.g., an anticodon loop, the acceptor stem, D arm or loop, variable loop, TψC arm or loop, other regions of the tRNA molecule, or a combination thereof. Typically, mutations in a leucyl tRNA include introducing a $CU(X)_n$ XXXAA sequence into the anticodon loop, where X refers to any nucleotide, and $(X)_n$ is optionally present. The n refers to number of bases the anticodon loop needs to be extended based on the selector codon, e.g., an extended codon, such as a four-, five-, six- base pair, etc. In one embodiment, mutations include matched base pairs in the stem region. In one embodiment, mutations include a first pair selected from T28:A42, G28:C42; C28:G42, etc. and a second pair selected from G49:C65 or C49:G65. The numbering refers to the positions on a tRNA molecule, e.g., see **Figure 4, Panel A**. The method can further include adding an additional sequence (CCA) to 3' terminus of the O-tRNA and/or measuring suppression activity.

[0082] Typically, an O-tRNA is obtained by subjecting to negative selection a population of cells of a first species, where the cells comprise a member of the plurality of potential O-tRNAs. The negative selection eliminates cells that comprise a member of the plurality of potential O-tRNAs that is aminoacylated by an aminoacyl-tRNA synthetase (RS) that is endogenous to the cells. This provides a pool of tRNAs that are orthogonal to the cell of the first species.

[0083] In certain embodiment in the negative selection, a selector codon(s) is introduced into polynucleotide that encodes a negative selection marker, e.g., an enzyme that confers antibiotic resistance, e.g., β-lactamase, an enzyme that confers a detectable product, e.g., β-galactosidase, chloramphenicol acetyltransferase (CAT), e.g., a toxic product, such as barnase, at a nonessential position, etc. Screening/selection can be done by growing the population of cells in the presence of a selective agent (e.g., an antibiotic, such as ampicillin). In one embodiment, the concentration of the selection agent is varied.

[0084] For example, to measure the activity of suppressor leucyl tRNAs, a selection system is used that is based on the *in vivo* suppression of selector codon, e.g., nonsense or frameshift mutations introduced into a polynucleotide that encodes a negative selection marker, e.g., a gene for β-lactamase (*bla*). For example, polynucleotide variants, e.g., *bla* variants, with, e.g., TAG, AGGA, and TGA, at a certain position (e.g., A184), are constructed. Cells, e.g., bacteria, are transformed with these polynucleotides. In the case of an orthogonal leucyl tRNA, which cannot be efficiently charged by endogenous *E. coli* synthetases, antibiotic resistance, e.g., ampicillin resistance, should be about or less than that for a bacteria transformed with no plasmid. If the leucyl tRNA is not orthogonal, or if a heterologous synthetase capable of charging the tRNA is co-expressed in the system, a higher level of antibiotic, e.g., ampicillin, resistance is be observed. Cells, e.g., bacteria, are chosen that are unable to grow on LB agar plates with antibiotic concentrations about equal to cells transformed with no plasmids.

[0085] In the case of a toxic product (e.g., ribonuclease or barnase), when a member of the plurality of potential leucyl tRNAs is aminoacylated by endogenous host, e.g., *Escherichia coli* synthetases (i.e., it is not orthogonal to the host, e.g., *Escherichia coli* synthetases), the selector codon is suppressed and the toxic polynucleotide product

produced leads to cell death. Cells harboring orthogonal leucyl tRNAs or non-functional tRNAs survive.

[0086]   In one embodiment, the pool of tRNAs that are orthogonal to a desired organism are then subjected to a positive selection in which a selector codon is placed in a positive selection marker, e.g., encoded by a drug resistance gene, such a β-lactamase gene. The positive selection is performed on cell comprising a polynucleotide encoding or comprising a member of the pool of tRNAs, a polynucleotide encoding a positive selection marker, and a polynucleotide encoding a cognate RS. These polynucleotides are expressed in the cell and the cell is grown in the presence of a selection agent, e.g., ampicillin. Leucyl tRNAs are then selected for their ability to be aminoacylated by the coexpressed cognate synthetase and to insert an amino acid in response to this selector codon. Typically, these cells show an enhancement in suppression efficiency compared to cells harboring non-functional tRNA(s), or tRNAs that cannot efficiently be recognized by the synthetase of interest. The cell harboring the non-functional tRNAs or tRNAs that are not efficiently recognized by the synthetase of interest, are sensitive to the antibiotic. Therefore, leucyl tRNAs that: (i) are not substrates for endogenous host, e.g., *Escherichia coli*, synthetases; (ii) can be aminoacylated by the synthetase of interest; and (iii) are functional in translation, survive both selections.

[0087]   The stringency of the selection, e.g., the positive selection, the negative selection or both the positive and negative selection, in the above described-methods, optionally include varying the selection stringency. For example, because barnase is an extremely toxic protein, the stringency of the negative selection can be controlled by introducing different numbers of selector codons into the barnase gene and/or by using an inducible promoter. In another example, the concentration of the selection or screening agent is varied (e.g., ampicillin concentration). In one aspect of the invention, the stringency is varied because the desired activity can be low during early rounds. Thus, less stringent selection criteria are applied in early rounds and more stringent criteria are applied in later rounds of selection. In certain embodiments, the negative selection, the positive selection or both the negative and positive selection, can be repeated multiple times. Multiple different negative selection markers, positive selection markers or both negative and positive selection markers, can be used. In certain embodiments, the positive and negative selection marker can be the same.

[0088]   Other types of selections/screening can be used in the invention for
producing orthogonal translational components, e.g., a leucyl O-tRNA, a leucyl O-RS, and a
leucyl O-tRNA/O-RS pair. For example, the negative selection marker, the positive
selection marker or both the positive and negative selection markers can include a marker

5   that fluoresces or catalyzes a luminescent reaction in the presence of a suitable reactant. In
another embodiment, a product of the marker is detected by fluorescence-activated cell
sorting (FACS) or by luminescence. Optionally, the marker includes an affinity based
screening marker. *See,* Francisco, J. A., et al., (1993) *Production and fluorescence-activated
cell sorting of Escherichia coli expressing a functional antibody fragment on the external*

10   *surface.* Proc Natl Acad Sci U S A. 90:10444-8.

[0089]   Additional general methods for producing a recombinant orthogonal tRNA
can be found, e.g., in International patent applications WO 2002/086075, entitled "Methods
and compositions for the production of orthogonal tRNA-aminoacyltRNA synthetase pairs;"
and, International Application Number PCT/2004/011786, filed April 16, 2004, entitled

15   "EXPANDING THE EUKARYOTIC GENETIC CODE." *See also,* Forster et al., (2003)
Programming peptidomimetic synthetases by translating genetic codes designed de novo
PNAS 100(11):6353-6357; and, Feng et al., (2003), *Expanding tRNA recognition of a tRNA
synthetase by a single amino acid change,* PNAS 100(10): 5676-5681. These are applied to
the present invention, e.g., using the substrates (e.g., leucyl-O-tRNAs or O-RSs) in such

20   available selection methods.

Orthogonal aminoacyl-tRNA synthetase (O-RS)
[0090]   A leucyl O-RS of the invention preferentially aminoacylates a leucyl O-
tRNA with a selected amino acid in vitro or in vivo. A leucyl O-RS of the invention can be
provided to the translation system, e.g., a cell, by a polypeptide that includes a leucyl O-RS

25   and/or by a polynucleotide that encodes a leucyl O-RS or a portion thereof. For example, a
leucyl O-RS, or a portion thereof, is encoded by a polynucleotide sequence as set forth in
any one of SEQ ID NO.: 13-14, or a complementary polynucleotide sequence thereof. In
another example, a leucyl O-RS comprises an amino acid sequence as set forth in any one of
SEQ ID NO.: 15-16, or a conservative variation thereof. *See, e.g.,* Table 3 and Example 2

30   herein for sequences of exemplary leucyl O-RS molecules.

[0091]   Methods for identifying an orthogonal aminoacyl-tRNA synthetase (O-RS),
e.g., a leucyl O-RS, for use with an O-tRNA, e.g., a leucyl O-tRNA, are also a feature of the

invention. For example, a method includes subjecting to positive selection a population of cells of a first species, where the cells individually comprise: 1) a member of a plurality of aminoacyl-tRNA synthetases (RSs), where the plurality of RSs comprise mutant RSs, RSs derived from a species other than the first species or both mutant RSs and RSs derived from

5   a species other than the first species; 2) the orthogonal tRNA (O-tRNA) from a second species; and 3) a polynucleotide that encodes a positive selection marker and comprises at least one selector codon. Cells are selected or screened for those that show an enhancement in suppression efficiency compared to cells lacking or with a reduced amount of the member of the plurality of RSs. Cells having an enhancement in suppression efficiency

10   comprise an active RS that aminoacylates the O-tRNA. A level of aminoacylation (in vitro or in vivo) by the active RS of a first set of tRNAs from the first species is compared to the level of aminoacylation (in vitro or in vivo) by the active RS of a second set of tRNAs from the second species. The level of aminoacylation can be determined by a detectable substance (e.g., a labeled amino acid or unnatural amino acid). The active RS that more

15   efficiently aminoacylates the second set of tRNAs compared to the first set of tRNAs is selected, thereby providing an efficient (optimized) orthogonal aminoacyl-tRNA synthetase for use with the O-tRNA. An O-RS, e.g., a leucyl O-RS, identified by the method, is also a feature of the invention.

[0092]   Any of a number of assays can be used to determine aminoacylation. These
20   assays can be performed *in vitro* or *in vivo*. For example, in vitro aminoacylation assays are described in, e.g., Hoben, P., and Soll, D. (1985) Methods Enzymol. 113:55-59. Aminoacylation can also be determined by using a reporter along with orthogonal translation components and detecting the reporter in a cell expressing a polynucleotide comprising at least one selector codon that encodes a protein. *See also*, WO 2002/085923,
25   entitled "IN VIVO INCORPORATION OF UNNATURAL AMINO ACIDS;" and, USSN 60/479,931 entitled "EXPANDING THE EUKARYOTIC GENETIC CODE."

[0093]   Identified leucyl O-RS can be further manipulated to alter the substrate specificity of the synthetase, so that only a desired unnatural amino acid, but not any of the common 20 amino acids are charged to the leucyl O-tRNA. Methods to generate an
30   orthogonal leucyl aminoacyl tRNA synthetase with a substrate specificity for an unnatural amino acid include mutating the synthetase, e.g., at the active site in the synthetase, at the editing mechanism site in the synthetase, at different sites by combining different domains

of synthetases, or the like, and applying a selection process. A strategy is used, which is based on the combination of a positive selection followed by a negative selection. In the positive selection, suppression of the selector codon introduced at a nonessential position(s) of a positive marker allows cells to survive under positive selection pressure. In the presence of both natural and unnatural amino acids, survivors thus encode active synthetases charging the orthogonal suppressor tRNA with either a natural or unnatural amino acid. In the negative selection, suppression of a selector codon introduced at a nonessential position(s) of a negative marker removes synthetases with natural amino acid specificities. Survivors of the negative and positive selection encode synthetases that aminoacylate (charge) the orthogonal suppressor tRNA with unnatural amino acids only. These synthetases can then be subjected to further mutagenesis, e.g., DNA shuffling or other recursive mutagenesis methods.

[0094]   The library of mutant leucyl O-RSs can be generated using various mutagenesis techniques known in the art. For example, the mutant RSs can be generated by site-specific mutations, random point mutations, homologous recombination, DNA shuffling or other recursive mutagenesis methods, chimeric construction or any combination thereof. For example, a library of mutant leucyl RSs can be produced from two or more other, e.g., smaller, less diverse "sub-libraries." Chimeric libraries of RSs are also included in the invention. It should be noted that libraries of tRNA synthetases from various organism (e.g., microorganisms such as eubacteria or archaebacteria) such as libraries that comprise natural diversity (see, e.g., U.S. Patent No. 6,238,884 to Short et al; U.S. Patent No. 5,756,316 to Schallenberger et al; U.S. Patent No. 5,783,431 to Petersen et al; U.S. Patent No. 5,824,485 to Thompson et al; U.S. Patent No. 5,958,672 to Short et al), are optionally constructed and screened for orthogonal pairs.

[0095]   Once the synthetases are subject to the positive and negative selection/screening strategy, these synthetases can then be subjected to further mutagenesis. For example, a nucleic acid that encodes the leucyl O-RS can be isolated; a set of polynucleotides that encode mutated leucyl O-RSs (e.g., by random mutagenesis, site-specific mutagenesis, recombination or any combination thereof) can be generated from the nucleic acid; and, these individual steps or a combination of these steps can be repeated until a mutated leucyl O-RS is obtained that preferentially aminoacylates the leucyl O-

tRNA with the unnatural amino acid. In one aspect of the invention, the steps are performed multiple times, e.g., at least two times.

[0096]   Additional levels of selection/screening stringency can also be used in the methods of the invention, for producing leucyl O-tRNA, leucyl O-RS, or pairs thereof. The selection or screening stringency can be varied on one or both steps of the method to produce an O-RS. This could include, e.g., varying the amount of selection/screening agent that is used, etc. Additional rounds of positive and/or negative selections can also be performed. Selecting or screening can also comprise one or more positive or negative selection or screening that includes, e.g., a change in amino acid permeability, a change in translation efficiency, a change in translational fidelity, etc. Typically, the one or more change is based upon a mutation in one or more gene in an organism in which an orthogonal tRNA-tRNA synthetase pair is used to produce protein.

[0097]   Additional general details for producing O-RS, and altering the substrate specificity of the synthetase can be found in WO 2002/086075 entitled "Methods and compositions for the production of orthogonal tRNA-aminoacyltRNA synthetase pairs; " and International Application Number PCT/US2004/011786, filed April 16, 2004.

SOURCE AND HOST ORGANISMS

[0098]   The translational components of the invention can be derived from non-eukaryotic organisms. For example, the orthogonal O-tRNA can be derived from a non-eukaryotic organism, e.g., an archaebacterium, such as *Methanococcus jannaschii, Methanobacterium thermoautotrophicum, Halobacterium* such as *Haloferax volcanii* and *Halobacterium* species *NRC-1, Archaeoglobus fulgidus, Pyrococcus furiosus, Pyrococcus horikoshii, Aeuropyrum pernix,* or the like, or a eubacterium, such as *Escherichia coli, Thermus thermophilus, Bacillus stearothermphilus,* or the like, while the orthogonal O-RS can be derived from a non-eukaryotic organism, e.g., *Methanobacterium thermoautotrophicum, Halobacterium* such as *Haloferax volcanii* and *Halobacterium* species *NRC-1, Archaeoglobus fulgidus, Pyrococcus furiosus, Pyrococcus horikoshii, Aeuropyrum pernix,* or the like, or a eubacterium, such as *Escherichia coli, Thermus thermophilus, Bacillus stearothermphilus,* or the like. In one embodiment, eukaryotic sources can also be used, e.g., plants, algae, protists, fungi, yeasts, animals (e.g., mammals, insects, arthropods, etc.), or the like.

[0099]   The individual components of a leucyl O-tRNA/O-RS pair can be derived from the same organism or different organisms. In one embodiment, the leucyl O-tRNA/O-RS pair is from the same organism. Alternatively, the leucyl O-tRNA and the leucyl O-RS of the leucyl O-tRNA/O-RS pair are from different organisms. For example, the leucyl O-tRNA can be derived from, e.g., a *Halobacterium sp NRC-1*, and the leucyl O-RS can be derived from, e.g., a *Methanobacterium thermoautrophicum*.

[0100]   The leucyl O-tRNA, leucyl O-RS or leucyl O-tRNA/O-RS pair can be selected or screened in vivo or in vitro and/or used in a cell, e.g., a non-eukaryotic cells (such as *E. coli* cell), or a eukaryotic cell, to produce a polypeptide with a selected amino acid (e.g., an unnatural amino acid). A non-eukaryotic cell can be from a variety of sources, e.g., *Methanobacterium thermoautotrophicum*, *Halobacterium* such as *Haloferax volcanii* and *Halobacterium* species *NRC-1*, *Archaeoglobus fulgidus*, *Pyrococcus furiosus*, *Pyrococcus horikoshii*, *Aeuropyrum pernix*, or the like, or a eubacterium, such as *Escherichia coli*, *Thermus thermophilus*, *Bacillus stearothermphilus*, or the like. A eukaryotic cell can be from any of a variety of sources, e.g., a plant (e.g., complex plant such as monocots, or dicots), an algae, a protist, a fungus, a yeast (e.g., *Saccharomyces cerevisiae*), an animal (e.g., a mammal, an insect, an arthropod, etc.), or the like. Compositions of cells with translational components of the invention are also a feature of the invention.

[0101]   *See also*, International Application Number PCT/US2004/011786, filed April 16, 2004, entitled "Expanding the Eukaryotic Genetic Code" for screening O-tRNA and/or O-RS in one species for use in another species.

SELECTOR CODONS

[0102]   Selector codons of the invention expand the genetic codon framework of protein biosynthetic machinery. For example, a selector codon includes, e.g., a unique three base codon, a nonsense codon, such as a stop codon, e.g., an amber codon (UAG), or an opal codon (UGA), an unnatural codon, at least a four base codon, a rare codon, or the like. A number of selector codons can be introduced into a desired gene, e.g., one or more, two or more, more than three, etc.

[0103]   In one embodiment, the methods involve the use of a selector codon that is a stop codon for the incorporation of a selected amino acid, e.g., an unnatural amino acids, in

vivo in a cell. For example, a leucyl O-tRNA is produced that recognizes the stop codon and is aminoacylated by a leucyl O-RS with a selected amino acid. This leucyl O-tRNA is not recognized by the naturally occurring host's aminoacyl-tRNA synthetases. Conventional site-directed mutagenesis can be used to introduce the stop codon at the site of

5   interest in a polypeptide of interest. *See, e.g.,* Sayers, J.R., et al. (1988), *5',3' Exonuclease in phosphorothioate-based oligonucleotide-directed mutagenesis.* Nucleic Acids Res, 791-802. When the leucyl O-RS, leucyl O-tRNA and the nucleic acid that encodes a polypeptide of interest are combined, e.g., in vivo, the selected amino acid is incorporated in response to the stop codon to give a polypeptide containing the selected amino acid, e.g., an

10   unnatural amino acid, at the specified position. In one embodiment of the invention, a stop codon used as a selector codon is an amber codon, UAG, and/or an opal codon, UGA. For example, see SEQ ID NO: 3 for an example of a leucyl O-tRNA that recognizes an amber codon, and see SEQ ID NO: 7 for an example of a leucyl O-tRNA that recognizes an opal codon. A genetic code in which UAG and UGA are both used as a selector codon can

15   encode 22 amino acids while preserving the ochre nonsense codon, UAA, which is the most abundant termination signal.

[0104]   The incorporation of selected amino acids, e.g., unnatural amino acids, *in vivo,* can be done without significant perturbation of the host cell. For example, in non-eukaryotic cells, such as *Escherichia coli,* because the suppression efficiency for the UAG

20   codon depends upon the competition between the O-tRNA, e.g., the amber suppressor tRNA, and the release factor 1 (RF1) (which binds to the UAG codon and initiates release of the growing peptide from the ribosome), the suppression efficiency can be modulated by, e.g., either increasing the expression level of O-tRNA, e.g., the suppressor tRNA, or using an RF1 deficient strain. In eukaryotic cells, because the suppression efficiency for the UAG

25   codon depends upon the competition between the O-tRNA, e.g., the amber suppressor tRNA, and a eukaryotic release factor (e.g., eRF) (which binds to a stop codon and initiates release of the growing peptide from the ribosome), the suppression efficiency can be modulated by, e.g., increasing the expression level of O-tRNA, e.g., the suppressor tRNA.

[0105]   Unnatural amino acids can also be encoded with rare codons. For example,

30   when the arginine concentration in an in vitro protein synthesis reaction is reduced, the rare arginine codon, AGG, has proven to be efficient for insertion of Ala by a synthetic tRNA acylated with alanine. *See, e.g.,* Ma et al., Biochemistry, 32:7939 (1993). In this case, the

synthetic tRNA competes with the naturally occurring tRNAArg, which exists as a minor species in *Escherichia coli*. Some organisms do not use all triplet codons. An unassigned codon AGA in *Micrococcus luteus* has been utilized for insertion of amino acids in an in vitro transcription/translation extract. *See, e.g.*, Kowal and Oliver, Nucl. Acid. Res., 25:4685 (1997). Components of the present invention can be generated to use these rare codons in vivo.

[0106]   Selector codons can also comprise extended codons, e.g., four or more base codons, such as, four, five, six or more base codons. Examples of four base codons include, e.g., AGGA, CUAG, UAGA, CCCU, and the like. Examples of five base codons include, e.g., AGGAC, CCCCU, CCCUC, CUAGA, CUACU, UAGGC, and the like. Methods of the invention include using extended codons based on frameshift suppression. Four or more base codons can insert, e.g., one or multiple selected amino acids, e.g., unnatural amino acids, into the same protein. For example, in the presence of mutated leucyl O-tRNAs, e.g., a special frameshift suppressor tRNAs, with anticodon loops, e.g., with a $CU(X)_n$ XXXAA sequence (where n=1), the four or more base codon is read as single amino acid. For example, see SEQ ID NOs.: 6 and 12 for leucyl O-tRNAs that recognize a four base codon. In other embodiments, the anticodon loops can decode, e.g., at least a four-base codon, at least a five-base codon, or at least a six-base codon or more. Since there are 256 possible four-base codons, multiple unnatural amino acids can be encoded in the same cell using a four or more base codon. *See also*, Anderson et al., (2002) *Exploring the Limits of Codon and Anticodon Size*, Chemistry and Biology, 9:237-244; Magliery, (2001) *Expanding the Genetic Code: Selection of Efficient Suppressors of Four-base Codons and Identification of "Shifty" Four-base Codons with a Library Approach in Escherichia coli*, J. Mol. Biol. 307: 755-769.

[0107]   For example, four-base codons have been used to incorporate unnatural amino acids into proteins using in vitro biosynthetic methods. *See, e.g.*, Ma et al., (1993) Biochemistry, 32:7939; and Hohsaka et al., (1999) J. Am. Chem. Soc., 121:34. CGGG and AGGU were used to simultaneously incorporate 2-naphthylalanine and an NBD derivative of lysine into streptavidin in vitro with two chemically acylated frameshift suppressor tRNAs. *See, e.g.*, Hohsaka et al., (1999) J. Am. Chem. Soc., 121:12194. In an in vivo study, Moore et al. examined the ability of tRNALeu derivatives with NCUA anticodons to suppress UAGN codons (N can be U, A, G, or C), and found that the quadruplet UAGA can

be decoded by a tRNALeu with a UCUA anticodon with an efficiency of 13 to 26% with

little decoding in the 0 or −1 frame. *See*, Moore et al., (2000) J. Mol. Biol., 298:195. In one

embodiment, extended codons based on rare codons or nonsense codons can be used in

invention, which can reduce missense readthrough and frameshift suppression at other

5   unwanted sites.

[0108]   For a given system, a selector codon can also include one of the natural three

base codons, where the endogenous system does not use (or rarely uses) the natural base

codon. For example, this includes a system that is lacking a tRNA that recognizes the

natural three base codon, and/or a system where the three base codon is a rare codon.

10   [0109]   Selector codons optionally include unnatural base pairs. These unnatural

base pairs further expand the existing genetic alphabet. One extra base pair increases the

number of triplet codons from 64 to 125. Properties of third base pairs include stable and

selective base pairing, efficient enzymatic incorporation into DNA with high fidelity by a

polymerase, and the efficient continued primer extension after synthesis of the nascent

15   unnatural base pair. Descriptions of unnatural base pairs which can be adapted for methods

and compositions include, e.g., Hirao, et al., (2002) *An unnatural base pair for*

incorporating amino acid analogues into protein, Nature Biotechnology, 20:177-182. See,

also, Wu, Y., et al., (2002) J. Am. Chem. Soc. 124:14626-14630. Other relevant

publications are listed herein.

20   [0110]   For in vivo usage, the unnatural nucleoside is membrane permeable and is

phosphorylated to form the corresponding triphosphate. In addition, the increased genetic

information is stable and not destroyed by cellular enzymes. Previous efforts by Benner and

others took advantage of hydrogen bonding patterns that are different from those in

canonical Watson-Crick pairs, the most noteworthy example of which is the iso-C:iso-G

25   pair. *See, e.g.,* Switzer et al., (1989) J. Am. Chem. Soc., 111:8322; and Piccirilli et al.,

(1990) Nature, 343:33; Kool, (2000) Curr. Opin. Chem. Biol., 4:602. These bases in

general mispair to some degree with natural bases and cannot be enzymatically replicated.

Kool and co-workers demonstrated that hydrophobic packing interactions between bases

can replace hydrogen bonding to drive the formation of base pair. *See*, Kool, (2000) Curr.

30   Opin. Chem. Biol., 4:602; and Guckian and Kool, (1998) Angew. Chem. Int. Ed. Engl., 36,

2825. In an effort to develop an unnatural base pair satisfying all the above requirements,

Schultz, Romesberg and co-workers have systematically synthesized and studied a series of

unnatural hydrophobic bases. A PICS:PICS self-pair is found to be more stable than natural base pairs, and can be efficiently incorporated into DNA by Klenow fragment of Escherichia coli DNA polymerase I (KF). *See, e.g.,* McMinn et al., (1999) J. Am. Chem. Soc., 121:11586; and Ogawa et al., (2000) J. Am. Chem. Soc., 122:3274. A 3MN:3MN self-pair can be synthesized by KF with efficiency and selectivity sufficient for biological function. *See, e.g.,* Ogawa et al., (2000) J. Am. Chem. Soc., 122:8803. However, both bases act as a chain terminator for further replication. A mutant DNA polymerase has been recently evolved that can be used to replicate the PICS self pair. In addition, a 7AI self pair can be replicated. See, e.g., Tae et al., (2001) J. Am. Chem. Soc., 123:7439. A novel metallobase pair, Dipic:Py, has also been developed, which forms a stable pair upon binding Cu(II). See, Meggers et al., (2000) J. Am. Chem. Soc., 122:10714. Because extended codons and unnatural codons are intrinsically orthogonal to natural codons, the methods of the invention can take advantage of this property to generate orthogonal tRNAs for them.

[0111]   A translational bypassing system can also be used to incorporate a selected amino acid, e.g., an unnatural amino acid, in a desired polypeptide. In a translational bypassing system, a large sequence is inserted into a gene but is not translated into protein. The sequence contains a structure that serves as a cue to induce the ribosome to hop over the sequence and resume translation downstream of the insertion.

SELECTED AND UNNATURAL AMINO ACIDS

[0112]   As used herein, a selected amino acid refers to any desired naturally occurring amino acid or unnatural amino acid. A naturally occurring amino acid includes any one of the twenty genetically encoded alpha-amino acids: alanine, arginine, asparagine, aspartic acid, cysteine, glutamine, glutamic acid, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, proline, serine, threonine, tryptophan, tyrosine, valine. In one embodiment, the selected amino acid is incorporated into a growing polypeptide chain with high fidelity, e.g., at greater than 75% efficiency for a given selector codon, at greater than about 80% efficiency for a given selector codon, at greater than about 90% efficiency for a given selector codon, at greater than about 95% efficiency for a given selector codon, or at greater than about 99% or more efficiency for a given selector codon.

[0113]   As used herein, an unnatural amino acid refers to any amino acid, modified amino acid, or amino acid analogue other than selenocysteine and/or pyrrolysine and the

following twenty genetically encoded alpha-amino acids: alanine, arginine, asparagine, aspartic acid, cysteine, glutamine, glutamic acid, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, proline, serine, threonine, tryptophan, tyrosine, valine. The generic structure of an alpha-amino acid is illustrated by Formula I:

I

$$H_2N \overset{R}{\wedge} CO_2H$$

5
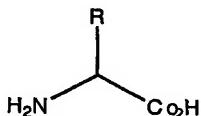
[0114]   An unnatural amino acid is typically any structure having Formula I wherein the R group is any substituent other than one used in the twenty natural amino acids. See, e.g., *Biochemistry* by L. Stryer, 3rd ed. 1988, Freeman and Company, New York, for

10   structures of the twenty natural amino acids. Note that, the unnatural amino acids of the invention can be naturally occurring compounds other than the twenty alpha-amino acids above.

[0115]   Because the unnatural amino acids of the invention typically differ from the natural amino acids in side chain only, the unnatural amino acids form amide bonds with

15   other amino acids, e.g., natural or unnatural, in the same manner in which they are formed in naturally occurring proteins. However, the unnatural amino acids have side chain groups that distinguish them from the natural amino acids.

[0116]   Because the unnatural amino acids of the invention typically differ from the natural amino acids in side chain, the unnatural amino acids form amide bonds with other

20   amino acids, e.g., natural or unnatural, in the same manner in which they are formed in naturally occurring proteins. However, the unnatural amino acids have side chain groups that distinguish them from the natural amino acids. For example, R in Formula I optionally comprises an alkyl-, aryl-, acyl-, keto-, azido-, hydroxyl-, hydrazine, cyano-, halo-, hydrazide, alkenyl, alkynl, ether, thiol, seleno-, sulfonyl-, borate, boronate, phospho,

25   phosphono, phosphine, heterocyclic, enone, imine, aldehyde, ester, thioacid, hydroxylamine, amine, and the like, or any combination thereof. In some embodiments, the unnatural amino acids have a photoactivatable cross-linker that is used, e.g., to link a

protein to a solid support. In one embodiment, the unnatural amino acids have a saccharide moiety attached to the amino acid side chain.

[0117] In addition to unnatural amino acids that contain novel side chains, unnatural amino acids also optionally comprise modified backbone structures, e.g., as illustrated by
5 the structures of Formula II and III:

II

R

Z—C—YH
‖
X

III

R   R'

H_2N CO_2H
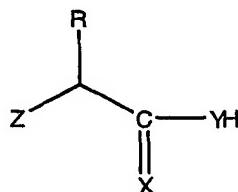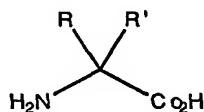
10 wherein Z typically comprises OH, NH_2, SH, NH-R', or S-R'; X and Y, which can be the same or different, typically comprise S or O, and R and R', which are optionally the same or different, are typically selected from the same list of constituents for the R group described above for the unnatural amino acids having Formula I as well as hydrogen. For example, unnatural amino acids of the invention optionally comprise substitutions in the amino or
15 carboxyl group as illustrated by Formulas II and III. Unnatural amino acids of this type include, but are not limited to, α-hydroxy acids, α-thioacids α-aminothiocarboxylates, e.g., with side chains corresponding to the common twenty natural amino acids or unnatural side chains. In addition, substitutions at the α-carbon optionally include L, D, or α-α-disubstituted amino acids such as D-glutamate, D-alanine, D-methyl-O-tyrosine,
20 aminobutyric acid, and the like. Other structural alternatives include cyclic amino acids, such as proline analogues as well as 3,4,6,7,8, and 9 membered ring proline analogues, β and γ amino acids such as substituted β-alanine and γ-amino butyric acid.

[0118]   For example, many unnatural amino acids are based on natural amino acids, such as tyrosine, glutamine, phenylalanine, and the like. Tyrosine analogs include para-substituted tyrosines, ortho-substituted tyrosines, and meta substituted tyrosines, wherein the substituted tyrosine comprises an acetyl group, a benzoyl group, an amino group, a hydrazine, an hydroxyamine, a thiol group, a carboxy group, an isopropyl group, a methyl group, a C_6 - C_{20} straight chain or branched hydrocarbon, a saturated or unsaturated hydrocarbon, an O-methyl group, a polyether group, a nitro group, or the like. In addition, multiply substituted aryl rings are also contemplated. Glutamine analogs of the invention include, but are not limited to, α-hydroxy derivatives, γ-substituted derivatives, cyclic derivatives, and amide substituted glutamine derivatives. Example phenylalanine analogs include, but are not limited to, para-substituted phenylalanines, ortho-substituted phenyalanines, and meta-substituted phenylalanines, wherein the substituent comprises a hydroxy group, a methoxy group, a methyl group, an allyl group, an aldehyde or keto group, or the like. Specific examples of unnatural amino acids include, but are not limited to, a p-acetyl-L- phenylalanine, a p-propargyl-phenylalanine, O-methyl-L-tyrosine, an L-3-(2-naphthyl)alanine, a 3-methyl-phenylalanine, an O-4-allyl-L-tyrosine, a 4-propyl-L-tyrosine, a tri-O-acetyl-GlcNAcβ-serine, an L-Dopa, a fluorinated phenylalanine, an isopropyl-L-phenylalanine, a p-azido-L-phenylalanine, a p-acyl-L-phenylalanine, a p-benzoyl-L-phenylalanine, an L-phosphoserine, a phosphonoserine, a phosphonotyrosine, a p-iodo-phenylalanine, a p-bromophenylalanine, a p-amino-L-phenylalanine, and an isopropyl-L-phenylalanine, and the like. The structures of a variety of unnatural amino acids are provided in, for example, Figures 16, 17, 18, 19, 26, and 29 of WO 2002/085923 entitled "In vivo incorporation of unnatural amino acids."

Chemical Synthesis of Unnatural Amino Acids

[0119]   Many of the unnatural amino acids provided above are commercially available, e.g., from Sigma (USA) or Aldrich (Milwaukee, WI, USA). Those that are not commercially available are optionally synthesized as provided in various publications or using standard methods known to those of skill in the art. For organic synthesis techniques, see, e.g., Organic Chemistry by Fessendon and Fessendon, (1982, Second Edition, Willard Grant Press, Boston Mass.); Advanced Organic Chemistry by March (Third Edition, 1985, Wiley and Sons, New York); and Advanced Organic Chemistry by Carey and Sundberg (Third Edition, Parts A and B, 1990, Plenum Press, New York). Additional publications

describing the synthesis of unnatural amino acids include, e.g., WO 2002/085923 entitled

"In vivo incorporation of Unnatural Amino Acids;" Matsoukas et al., (1995) *J. Med. Chem.*,

38, 4660-4669; King, F.E. & Kidd, D.A.A. (1949) *A New Synthesis of Glutamine and of γ-*

Dipeptides of Glutamic Acid from Phthylated Intermediates. J. Chem. Soc., 3315-3319;

5   Friedman, O.M. & Chatterrji, R. (1959) *Synthesis of Derivatives of Glutamine as Model*

Substrates for Anti-Tumor Agents. J. Am. Chem. Soc. 81, 3750-3752; Craig, J.C. et al.

(1988) *Absolute Configuration of the Enantiomers of 7-Chloro-4 [[4-(diethylamino)-1-*

methylbutyl]amino]quinoline (Chloroquine). J. Org. Chem. 53, 1167-1170; Azoulay, M.,

Vilmont, M. & Frappier, F. (1991) *Glutamine analogues as Potential Antimalarials,. Eur. J.*

10   *Med. Chem.* 26, 201-5; Koskinen, A.M.P. & Rapoport, H. (1989) *Synthesis of 4-Substituted*

Prolines as Conformationally Constrained Amino Acid Analogues. J. Org. Chem. 54, 1859-

1866; Christie, B.D. & Rapoport, H. (1985) *Synthesis of Optically Pure Pipecolates from L-*

Asparagine. Application to the Total Synthesis of (+)-Apovincamine through Amino Acid

Decarbonylation and Iminium Ion Cyclization. J. Org. Chem. 1989:1859-1866; Barton et

15   al., (1987) *Synthesis of Novel a-Amino-Acids and Derivatives Using Radical Chemistry:*

Synthesis of L- and D-a-Amino-Adipic Acids, L-a-aminopimelic Acid and Appropriate

Unsaturated Derivatives. Tetrahedron Lett. 43:4297-4308; and, Subasinghe et al., (1992)

Quisqualic acid analogues: synthesis of beta-heterocyclic 2-aminopropanoic acid

derivatives and their activity at a novel quisqualate-sensitized site. J. Med. Chem. 35:4602-

20   7. *See also*, International Application Number PCT/US03/41346, entitled "Protein Arrays,"

filed on December 22, 2003.

Cellular uptake of unnatural amino acids

[0120]   Unnatural amino acid uptake by a cell is one issue that is typically

considered when designing and selecting unnatural amino acids, e.g., for incorporation into

25   a protein. For example, the high charge density of α-amino acids suggests that these

compounds are unlikely to be cell permeable. Natural amino acids are taken up into the cell

via a collection of protein-based transport systems often displaying varying degrees of

amino acid specificity. A rapid screen can be done which assesses which unnatural amino

acids, if any, are taken up by cells. *See, e.g.,* the toxicity assays in, e.g., International

30   Application Number PCT/US03/41346, entitled "Protein Arrays," filed on December 22,

2003; and Liu, D.R. & Schultz, P.G. (1999) *Progress toward the evolution of an organism*

with an expanded genetic code. PNAS United States 96:4780-4785. Although uptake is

easily analyzed with various assays, an alternative to designing unnatural amino acids that are amenable to cellular uptake pathways is to provide biosynthetic pathways to create amino acids *in vivo*.

Biosynthesis of Unnatural Amino Acids

[0121]   Many biosynthetic pathways already exist in cells for the production of amino acids and other compounds. While a biosynthetic method for a particular unnatural amino acid may not exist in nature, e.g., in a cell, the invention provides such methods. For example, biosynthetic pathways for unnatural amino acids are optionally generated in host cell by adding new enzymes or modifying existing host cell pathways. Additional new enzymes are optionally naturally occurring enzymes or artificially evolved enzymes. For example, the biosynthesis of *p*-aminophenylalanine (as presented in an example in WO 2002/085923, *supra*) relies on the addition of a combination of known enzymes from other organisms. The genes for these enzymes can be introduced into a cell by transforming the cell with a plasmid comprising the genes. The genes, when expressed in the cell, provide an enzymatic pathway to synthesize the desired compound. Examples of the types of enzymes that are optionally added are provided in the examples below. Additional enzymes sequences are found, e.g., in Genbank. Artificially evolved enzymes are also optionally added into a cell in the same manner. In this manner, the cellular machinery and resources of a cell are manipulated to produce unnatural amino acids.

[0122]   Indeed, any of a variety of methods can be used for producing novel enzymes for use in biosynthetic pathways, or for evolution of existing pathways, for the production of unnatural amino acids, in vitro or in vivo. Many available methods of evolving enzymes and other biosynthetic pathway components can be applied to the present invention to produce unnatural amino acids (or, indeed, to evolve synthetases to have new substrate specificities or other activities of interest). For example, DNA shuffling is optionally used to develop novel enzymes and/or pathways of such enzymes for the production of unnatural amino acids (or production of new synthetases), in vitro or in vivo. *See, e.g.,* Stemmer (1994), *Rapid evolution of a protein in vitro by DNA shuffling*, Nature 370(4):389-391; and, Stemmer, (1994), *DNA shuffling by random fragmentation and reassembly: In vitro recombination for molecular evolution*, Proc. Natl. Acad. Sci. USA., 91:10747-10751. A related approach shuffles families of related (e.g., homologous) genes to quickly evolve enzymes with desired characteristics. An example of such "family gene

shuffling" methods is found in Crameri et al. (1998) "DNA shuffling of a family of genes from diverse species accelerates directed evolution" Nature, 391(6664): 288-291. New enzymes (whether biosynthetic pathway components or synthetases) can also be generated using a DNA recombination procedure known as "incremental truncation for the creation of

5   hybrid enzymes" ("ITCHY"), e.g., as described in Ostermeier et al. (1999) "A combinatorial approach to hybrid enzymes independent of DNA homology" Nature Biotech 17:1205. This approach can also be used to generate a library of enzyme or other pathway variants which can serve as substrates for one or more in vitro or in vivo recombination methods. See, also, Ostermeier et al. (1999) "Combinatorial Protein Engineering by

10   Incremental Truncation," Proc. Natl. Acad. Sci. USA, 96: 3562-67, and Ostermeier et al. (1999), "Incremental Truncation as a Strategy in the Engineering of Novel Biocatalysts," Biological and Medicinal Chemistry, 7: 2139-44. Another approach uses exponential ensemble mutagenesis to produce libraries of enzyme or other pathway variants that are, e.g., selected for an ability to catalyze a biosynthetic reaction relevant to producing an

15   unnatural amino acid (or a new synthetase). In this approach, small groups of residues in a sequence of interest are randomized in parallel to identify, at each altered position, amino acids which lead to functional proteins. Examples of such procedures, which can be adapted to the present invention to produce new enzymes for the production of unnatural amino acids (or new synthetases) are found in Delegrave & Youvan (1993) Biotechnology

20   Research 11:1548-1552. In yet another approach, random or semi-random mutagenesis using doped or degenerate oligonucleotides for enzyme and/or pathway component engineering can be used, e.g., by using the general mutagenesis methods of e.g., Arkin and Youvan (1992) "Optimizing nucleotide mixtures to encode specific subsets of amino acids for semi-random mutagenesis" Biotechnology 10:297-300; or Reidhaar-Olson et al. (1991)

25   "Random mutagenesis of protein sequences using oligonucleotide cassettes" Methods Enzymol. 208:564-86. Yet another approach, often termed a "non-stochastic" mutagenesis, which uses polynucleotide reassembly and site-saturation mutagenesis can be used to produce enzymes and/or pathway components, which can then be screened for an ability to perform one or more synthetase or biosynthetic pathway function (e.g., for the production of

30   unnatural amino acids in vivo). See, e.g., Short "Non-Stochastic Generation of Genetic Vaccines and Enzymes" WO 00/46344.

[0123]   An alternative to such mutational methods involves recombining entire genomes of organisms and selecting resulting progeny for particular pathway functions (often referred to as "whole genome shuffling"). This approach can be applied to the present invention, e.g., by genomic recombination and selection of an organism (e.g., an *E. coli* or other cell) for an ability to produce an unnatural amino acid (or intermediate thereof). For example, methods taught in the following publications can be applied to pathway design for the evolution of existing and/or new pathways in cells to produce unnatural amino acids in vivo: Patnaik et al. (2002) "Genome shuffling of lactobacillus for improved acid tolerance" Nature Biotechnology, 20(7): 707-712; and Zhang et al. (2002) "Genome shuffling leads to rapid phenotypic improvement in bacteria" Nature, February 7, 415(6872): 644-646.

[0124]   Other techniques for organism and metabolic pathway engineering, e.g., for the production of desired compounds are also available and can also be applied to the production of unnatural amino acids. Examples of publications teaching useful pathway engineering approaches include: Nakamura and White (2003) "Metabolic engineering for the microbial production of 1,3 propanediol" Curr. Opin. Biotechnol. 14(5):454-9; Berry et al. (2002) "Application of Metabolic Engineering to improve both the production and use of Biotech Indigo" J. Industrial Microbiology and Biotechnology 28:127-133; Banta et al. (2002) "Optimizing an artificial metabolic pathway: Engineering the cofactor specificity of Corynebacterium 2,5-diketo-D-gluconic acid reductase for use in vitamin C biosynthesis" Biochemistry, 41(20), 6226-36; Selivonova et al. (2001) "Rapid Evolution of Novel Traits in Microorganisms" Applied and Environmental Microbiology, 67:3645, and many others.

[0125]   Regardless of the method used, typically, the unnatural amino acid produced with an engineered biosynthetic pathway of the invention is produced in a concentration sufficient for efficient protein biosynthesis, e.g., a natural cellular amount, but not to such a degree as to significantly affect the concentration of other cellular amino acids or to exhaust cellular resources. Typical concentrations produced *in vivo* in this manner are about 10 mM to about 0.05 mM. Once a cell is engineered to produce enzymes desired for a specific pathway and an unnatural amino acid is generated, *in vivo* selections are optionally used to further optimize the production of the unnatural amino acid for both ribosomal protein synthesis and cell growth.

[0126]   As described above and below, the invention provides for nucleic acid polynucleotide sequences and polypeptide amino acid sequences, e.g., leucyl O-tRNAs and leucyl O-RSs, and, e.g., compositions, systems and methods comprising said sequences. Examples of said sequences, e.g., leucyl O-tRNAs and leucyl O-RSs are disclosed herein

5   (see, Table 3, e.g., SEQ ID NO. 1-7, 12-16). However, one of skill in the art will appreciate that the invention is not limited to those sequences disclosed herein, e.g., as in the Examples. One of skill will appreciate that the invention also provides many related and unrelated sequences with the functions described herein, e.g., encoding a leucyl O-tRNA or a leucyl O-RS.

10   [0127]   The invention provides polypeptides (leucyl O-RSs) and polynucleotides, e.g., leucyl O-tRNA, polynucleotides that encode leucyl O-RSs or portions thereof, oligonucleotides used to isolate aminoacyl-tRNA synthetase clones, etc. Polynucleotides of the invention include those that encode proteins or polypeptides of interest of the invention with one or more selector codon. In addition, polynucleotides of the invention include, e.g.,

15   a polynucleotide comprising a nucleotide sequence as set forth in any one of SEQ ID NO.: 1-2, 4-7 and 12; a polynucleotide that is complementary to or that encodes a polynucleotide sequence thereof. A polynucleotide of the invention also includes a polynucleotide that encodes a polypeptide of the invention. Similarly, a nucleic acid that hybridizes to a polynucleotide indicated above under highly stringent conditions over substantially the

20   entire length of the nucleic acid is a polynucleotide of the invention. In one embodiment, a composition includes a polypeptide of the invention and an excipient (e.g., buffer, water, pharmaceutically acceptable excipient, etc.). The invention also provides an antibody or antisera specifically immunoreactive with a polypeptide of the invention.

[0128]   A polynucleotide of the invention also includes a polynucleotide that is, e.g.,

25   at least 75%, at least 80%, at least 90%, at least 95%, at least 98% or more identical to that of a naturally occurring leucyl tRNA and comprises an anticodon loop comprising a CU(X)$_n$XXXAA sequence, an stem region lacking noncanonical base pairs and a conserved discriminator base at position 73. A polynucleotide also includes a polynucleotide that is , e.g., at least 75%, at least 80%, at least 90%, at least 95%, at least 98% or more identical to

30   that of a naturally occurring leucyl tRNA and comprises an anticodon loop comprising a CUUCCUAA sequence, a first pair selected from T28:A42, G28:C42 and/or C28:G42, and

a second pair selected from G:49:C65 or C49:G65, wherein the numbering corresponds to that indicated in **Figure 4, Panel A**.

[0129]   In certain embodiments, a vector (e.g., a plasmid, a cosmid, a phage, a virus, etc.) comprises a polynucleotide of the invention. In one embodiment, the vector is an

5   expression vector. In another embodiment, the expression vector includes a promoter operably linked to one or more of the polynucleotides of the invention. In another embodiment, a cell comprises a vector that includes a polynucleotide of the invention.

[0130]   One of skill will also appreciate that many variants of the disclosed sequences are included in the invention. For example, conservative variations of the

10   disclosed sequences that yield a functionally identical sequence are included in the invention. Variants of the nucleic acid polynucleotide sequences, wherein the variants hybridize to at least one disclosed sequence, are considered to be included in the invention. Unique subsequences of the sequences disclosed herein, as determined by, e.g., standard sequence comparison techniques, are also included in the invention.

15   Conservative variations
[0131]   Owing to the degeneracy of the genetic code, "silent substitutions" (*i.e.,* substitutions in a nucleic acid sequence which do not result in an alteration in an encoded polypeptide) are an implied feature of *every* nucleic acid sequence which encodes an amino acid. Similarly, "conservative amino acid substitutions," in one or a few amino acids in an

20   amino acid sequence are substituted with different amino acids with highly similar properties, are also readily identified as being highly similar to a disclosed construct. Such conservative variations of each disclosed sequence are a feature of the present invention.

[0132]   "Conservative variations" of a particular nucleic acid sequence refers to those nucleic acids which encode identical or essentially identical amino acid sequences, or,

25   where the nucleic acid does not encode an amino acid sequence, to essentially identical sequences. One of skill will recognize that individual substitutions, deletions or additions which alter, add or delete a single amino acid or a small percentage of amino acids (typically less than 5%, more typically less than 4%, 2% or 1%) in an encoded sequence are "conservatively modified variations" where the alterations result in the deletion of an amino

30   acid, addition of an amino acid, or substitution of an amino acid with a chemically similar amino acid. Thus, "conservative variations" of a listed polypeptide sequence of the present

invention include substitutions of a small percentage, typically less than 5%, more typically less than 2% or 1%, of the amino acids of the polypeptide sequence, with a conservatively selected amino acid of the same conservative substitution group. Finally, the addition of sequences which do not essentially alter the encoded activity of a nucleic acid molecule,

5   such as the addition of a non-functional sequence, is a conservative variation of the basic nucleic acid.

[0133]   Conservative substitution tables providing functionally similar amino acids are well known in the art. The following sets forth example groups which contain natural amino acids that include "conservative substitutions" for one another.

10

Conservative Substitution Groups

Nonpolar and/or Aliphatic Side Chains	Polar, Uncharged Side Chains	Aromatic Side Chains	Positively Charged Side Chains	Negatively Charged Side Chains
Glycine Alanine Valine Leucine Isoleucine Proline	Serine Threonine Cysteine Methionine Asparagine Glutamine	Phenylalanine Tyrosine Tryptophan	Lysine Arginine Histidine	Aspartate Glutamate

Nucleic Acid Hybridization

[0134]   Comparative hybridization can be used to identify nucleic acids of the invention, such as SEQ ID NO.: 1-2, 4-7 and 12, including conservative variations of

15 nucleic acids of the invention, and this comparative hybridization method is a preferred method of distinguishing nucleic acids of the invention. In addition, target nucleic acids which hybridize to the nucleic acids represented by SEQ ID NO: 1-2, 4-7 and 12 under high, ultra-high and ultra-ultra high stringency conditions are a feature of the invention. Examples of such nucleic acids include those with one or a few silent or conservative

20 nucleic acid substitutions as compared to a given nucleic acid sequence.

[0135]   A test nucleic acid is said to specifically hybridize to a probe nucleic acid when it hybridizes at least ½ as well to the probe as to the perfectly matched complementary target, i.e., with a signal to noise ratio at least ½ as high as hybridization of the probe to the target under conditions in which the perfectly matched probe binds to the perfectly matched

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complementary target with a signal to noise ratio that is at least about 5x-10x as high as that observed for hybridization to any of the unmatched target nucleic acids.

[0136]　　　　Nucleic acids "hybridize" when they associate, typically in solution. Nucleic acids hybridize due to a variety of well characterized physico-chemical forces, such as hydrogen bonding, solvent exclusion, base stacking and the like. An extensive guide to the hybridization of nucleic acids is found in Tijssen (1993) *Laboratory Techniques in Biochemistry and Molecular Biology--Hybridization with Nucleic Acid Probes* part I chapter 2, "Overview of principles of hybridization and the strategy of nucleic acid probe assays," (Elsevier, New York), as well as in Ausubel, *supra*. Hames and Higgins (1995) Gene Probes 1 IRL Press at Oxford University Press, Oxford, England, (Hames and Higgins 1) and Hames and Higgins (1995) Gene Probes 2 IRL Press at Oxford University Press, Oxford, England (Hames and Higgins 2) provide details on the synthesis, labeling, detection and quantification of DNA and RNA, including oligonucleotides.

[0137]　　　　An example of stringent hybridization conditions for hybridization of complementary nucleic acids which have more than 100 complementary residues on a filter in a Southern or northern blot is 50% formalin with 1 mg of heparin at 42°C, with the hybridization being carried out overnight. An example of stringent wash conditions is a 0.2x SSC wash at 65°C for 15 minutes (*see,* Sambrook, *supra* for a description of SSC buffer). Often the high stringency wash is preceded by a low stringency wash to remove background probe signal. An example low stringency wash is 2x SSC at 40°C for 15 minutes. In general, a signal to noise ratio of 5x (or higher) than that observed for an unrelated probe in the particular hybridization assay indicates detection of a specific hybridization.

[0138]　　　　"Stringent hybridization wash conditions" in the context of nucleic acid hybridization experiments such as Southern and northern hybridizations are sequence dependent, and are different under different environmental parameters. An extensive guide to the hybridization of nucleic acids is found in Tijssen (1993), *supra.* and in Hames and Higgins, 1 and 2. Stringent hybridization and wash conditions can easily be determined empirically for any test nucleic acid. For example, in determining highly stringent hybridization and wash conditions, the hybridization and wash conditions are gradually increased (e.g., by increasing temperature, decreasing salt concentration, increasing

detergent concentration and/or increasing the concentration of organic solvents such as formalin in the hybridization or wash), until a selected set of criteria are met. For example, the hybridization and wash conditions are gradually increased until a probe binds to a perfectly matched complementary target with a signal to noise ratio that is at least 5x as

5 high as that observed for hybridization of the probe to an unmatched target.

[0139]   "Very stringent" conditions are selected to be equal to the thermal melting point (T_m) for a particular probe. The T_m is the temperature (under defined ionic strength and pH) at which 50% of the test sequence hybridizes to a perfectly matched probe. For the purposes of the present invention, generally, "highly stringent" hybridization and wash

10 conditions are selected to be about 5° C lower than the T_m for the specific sequence at a defined ionic strength and pH.

[0140]   "Ultra high-stringency" hybridization and wash conditions are those in which the stringency of hybridization and wash conditions are increased until the signal to noise ratio for binding of the probe to the perfectly matched complementary target nucleic acid is

15 at least 10x as high as that observed for hybridization to any of the unmatched target nucleic acids. A target nucleic acid which hybridizes to a probe under such conditions, with a signal to noise ratio of at least ½ that of the perfectly matched complementary target nucleic acid is said to bind to the probe under ultra-high stringency conditions.

[0141]   Similarly, even higher levels of stringency can be determined by gradually

20 increasing the hybridization and/or wash conditions of the relevant hybridization assay. For example, those in which the stringency of hybridization and wash conditions are increased until the signal to noise ratio for binding of the probe to the perfectly matched complementary target nucleic acid is at least 10x, 20X, 50X, 100X, or 500X or more as high as that observed for hybridization to any of the unmatched target nucleic acids. A target

25 nucleic acid which hybridizes to a probe under such conditions, with a signal to noise ratio of at least ½ that of the perfectly matched complementary target nucleic acid is said to bind to the probe under ultra-ultra-high stringency conditions.

[0142]   Nucleic acids which do not hybridize to each other under stringent conditions are still substantially identical if the polypeptides which they encode are substantially

30 identical. This occurs, e.g., when a copy of a nucleic acid is created using the maximum codon degeneracy permitted by the genetic code.

Unique subsequences

[0143]   In one aspect, the invention provides a nucleic acid that comprises a unique subsequence in a nucleic acid selected from the sequences of leucyl O-tRNAs and leucyl O-RSs disclosed herein. The unique subsequence is unique as compared to a nucleic acid corresponding to any known leucyl O-tRNA or leucyl O-RS nucleic acid sequence. Alignment can be performed using, e.g., BLAST set to default parameters. Any unique subsequence is useful, e.g., as a probe to identify the nucleic acids of the invention.

[0144]   Similarly, the invention includes a polypeptide which comprises a unique subsequence in a polypeptide selected from the sequences of leucyl O-RSs disclosed herein. Here, the unique subsequence is unique as compared to a polypeptide corresponding to any of known polypeptide sequence.

[0145]   The invention also provides for target nucleic acids which hybridizes under stringent conditions to a unique coding oligonucleotide which encodes a unique subsequence in a polypeptide selected from the sequences of leucyl O-RSs wherein the unique subsequence is unique as compared to a polypeptide corresponding to any of the control polypeptides (e.g., parental sequences from which synthetases of the invention were derived, e.g., by mutation). Unique sequences are determined as noted above.

Sequence comparison, identity, and homology

[0146]   The terms "identical" or percent "identity," in the context of two or more nucleic acid or polypeptide sequences, refer to two or more sequences or subsequences that are the same or have a specified percentage of amino acid residues or nucleotides that are the same, when compared and aligned for maximum correspondence, as measured using one of the sequence comparison algorithms described below (or other algorithms available to persons of skill) or by visual inspection.

[0147]   The phrase "substantially identical," in the context of two nucleic acids or polypeptides (e.g., DNAs encoding an leucyl O-tRNA or leucyl O-RS, or the amino acid sequence of an O-RS) refers to two or more sequences or subsequences that have at least about 60%, about 80%, about 90-95%, about 98%, about 99% or more nucleotide or amino acid residue identity, when compared and aligned for maximum correspondence, as measured using a sequence comparison algorithm or by visual inspection. Such "substantially identical" sequences are typically considered to be "homologous," without reference to actual ancestry. Preferably, the "substantial identity" exists over a region of the

sequences that is at least about 50 residues in length, more preferably over a region of at least about 100 residues, and most preferably, the sequences are substantially identical over at least about 150 residues, or over the full length of the two sequences to be compared.

[0148]   For sequence comparison and homology determination, typically one sequence acts as a reference sequence to which test sequences are compared. When using a sequence comparison algorithm, test and reference sequences are input into a computer, subsequence coordinates are designated, if necessary, and sequence algorithm program parameters are designated. The sequence comparison algorithm then calculates the percent sequence identity for the test sequence(s) relative to the reference sequence, based on the designated program parameters.

[0149]   Optimal alignment of sequences for comparison can be conducted, *e.g.*, by the local homology algorithm of Smith & Waterman, Adv. Appl. Math. 2:482 (1981), by the homology alignment algorithm of Needleman & Wunsch, J. Mol. Biol. 48:443 (1970), by the search for similarity method of Pearson & Lipman, Proc. Nat'l. Acad. Sci. USA 85:2444 (1988), by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, WI), or by visual inspection (*see generally*, Ausubel *et al., infra*).

[0150]   One example of an algorithm that is suitable for determining percent sequence identity and sequence similarity is the BLAST algorithm, which is described in Altschul *et al.*, J. Mol. Biol. 215:403-410 (1990). Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information (www.ncbi.nlm.nih.gov/). This algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul *et al., supra*). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them. The word hits are then extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Cumulative scores are calculated using, for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always > 0) and N (penalty score for mismatching residues; always < 0). For amino acid sequences, a scoring matrix is used to

calculate the cumulative score. Extension of the word hits in each direction are halted

when: the cumulative alignment score falls off by the quantity X from its maximum

achieved value; the cumulative score goes to zero or below, due to the accumulation of one

or more negative-scoring residue alignments; or the end of either sequence is reached. The

5   BLAST algorithm parameters W, T, and X determine the sensitivity and speed of the

alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength

(W) of 11, an expectation (E) of 10, a cutoff of 100, M=5, N=-4, and a comparison of both

strands. For amino acid sequences, the BLASTP program uses as defaults a wordlength (W)

of 3, an expectation (E) of 10, and the BLOSUM62 scoring matrix (*see* Henikoff &

10   Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915).

[0151]   In addition to calculating percent sequence identity, the BLAST algorithm

also performs a statistical analysis of the similarity between two sequences (*see, e.g.*, Karlin

& Altschul, Proc. Nat'l. Acad. Sci. USA 90:5873-5787 (1993)). One measure of similarity

provided by the BLAST algorithm is the smallest sum probability (P(N)), which provides an

15   indication of the probability by which a match between two nucleotide or amino acid

sequences would occur by chance. For example, a nucleic acid is considered similar to a

reference sequence if the smallest sum probability in a comparison of the test nucleic acid to

the reference nucleic acid is less than about 0.1, more preferably less than about 0.01, and

most preferably less than about 0.001.

20   Mutagenesis and Other Molecular Biology Techniques
     [0152]   Polynucleotide and polypeptides of the invention and used in the invention

can be manipulated using molecular biological techniques. General texts which describe

molecular biological techniques include Berger and Kimmel, Guide to Molecular Cloning

Techniques, Methods in Enzymology volume 152 Academic Press, Inc., San Diego, CA

25   (Berger); Sambrook et al., Molecular Cloning - A Laboratory Manual (2nd Ed.), Vol. 1-3,

Cold Spring Harbor Laboratory, Cold Spring Harbor, New York, 1989 ("Sambrook") and

Current Protocols in Molecular Biology, F.M. Ausubel et al., eds., Current Protocols, a joint

venture between Greene Publishing Associates, Inc. and John Wiley & Sons, Inc.,

(supplemented through 1999) ("Ausubel")). These texts describe mutagenesis, the use of

30   vectors, promoters and many other relevant topics related to, e.g., the generation of genes

that include selector codons for production of proteins that include selected amino acids

(e.g., unnatural amino acids), leucyl orthogonal tRNAs, leucyl orthogonal synthetases, and pairs thereof.

[0153]   Various types of mutagenesis are used in the invention, e.g., to mutate tRNA molecules, to produce libraries of leucyl tRNAs, to produce libraries of leucyl synthetases, to insert selector codons that encode a selected amino acid in a protein or polypeptide of interest. They include but are not limited to site-directed, random point mutagenesis, homologous recombination, DNA shuffling or other recursive mutagenesis methods, chimeric construction, mutagenesis using uracil containing templates, oligonucleotide-directed mutagenesis, phosphorothioate-modified DNA mutagenesis, mutagenesis using gapped duplex DNA or the like, or any combination thereof. Additional suitable methods include point mismatch repair, mutagenesis using repair-deficient host strains, restriction-selection and restriction-purification, deletion mutagenesis, mutagenesis by total gene synthesis, double-strand break repair, and the like. Mutagenesis, e.g., involving chimeric constructs, is also included in the present invention. In one embodiment, mutagenesis can be guided by known information of the naturally occurring molecule or altered or mutated naturally occurring molecule, e.g., sequence, sequence comparisons, physical properties, crystal structure or the like.

[0154]   Host cells are genetically engineered (e.g., transformed, transduced or transfected) with the polynucleotides of the invention or constructs which include a polynucleotide of the invention, e.g., a vector of the invention, which can be, for example, a cloning vector or an expression vector. For example, the coding regions for the orthogonal tRNA, the orthogonal tRNA synthetase, and the protein to be derivatized are operably linked to gene expression control elements that are functional in the desired host cell. Typical vectors contain transcription and translation terminators, transcription and translation initiation sequences, and promoters useful for regulation of the expression of the particular target nucleic acid. The vectors optionally comprise generic expression cassettes containing at least one independent terminator sequence, sequences permitting replication of the cassette in eukaryotes, or prokaryotes, or both (e.g., shuttle vectors) and selection markers for both prokaryotic and eukaryotic systems. Vectors are suitable for replication and/or integration in prokaryotes, eukaryotes, or preferably both. *See,* Giliman & Smith, *Gene* 8:81 (1979); Roberts, *et al., Nature,* 328:731 (1987); Schneider, B., *et al., Protein Expr. Purif.* 6435:10 (1995); Ausubel, Sambrook, Berger (*all supra*). The vector can be, for

example, in the form of a plasmid, a bacterium, a virus, a naked polynucleotide, or a conjugated polynucleotide. The vectors are introduced into cells and/or microorganisms by standard methods including electroporation (From et al., Proc. Natl. Acad. Sci. USA 82, 5824 (1985), infection by viral vectors, high velocity ballistic penetration by small particles

5   with the nucleic acid either within the matrix of small beads or particles, or on the surface (Klein et al., Nature 327, 70-73 (1987)), and/or the like.

[0155]   A catalogue of Bacteria and Bacteriophages useful for cloning is provided, e.g., by the ATCC, e.g., The ATCC Catalogue of Bacteria and Bacteriophage (1992) Ghema et al. (eds) published by the ATCC. Additional basic procedures for sequencing,

10   cloning and other aspects of molecular biology and underlying theoretical considerations are also found in Watson et al. (1992) Recombinant DNA Second Edition Scientific American Books, NY. In addition, essentially any nucleic acid (and virtually any labeled nucleic acid, whether standard or non-standard) can be custom or standard ordered from any of a variety of commercial sources, such as the Midland Certified Reagent Company (Midland, TX

15   mcrc.com), The Great American Gene Company (Ramona, CA available on the World Wide Web at genco.com), ExpressGen Inc. (Chicago, IL available on the World Wide Web at expressgen.com), Operon Technologies Inc. (Alameda, CA) and many others.

[0156]   The engineered host cells can be cultured in conventional nutrient media modified as appropriate for such activities as, for example, screening steps, activating

20   promoters or selecting transformants. These cells can optionally be cultured into transgenic organisms. Other useful references, e.g. for cell isolation and culture (e.g., for subsequent nucleic acid isolation) include Freshney (1994) Culture of Animal Cells, a Manual of Basic Technique, third edition, Wiley- Liss, New York and the references cited therein; Payne et al. (1992) Plant Cell and Tissue Culture in Liquid Systems John Wiley & Sons, Inc. New

25   York, NY; Gamborg and Phillips (eds) (1995) Plant Cell, Tissue and Organ Culture; Fundamental Methods Springer Lab Manual, Springer-Verlag (Berlin Heidelberg New York) and Atlas and Parks (eds) The Handbook of Microbiological Media (1993) CRC Press, Boca Raton, FL.

PROTEINS AND POLYPEPTIDES OF INTEREST

30   [0157]   Methods of producing a protein in a cell with a selected amino acid at a specified position are also a feature of the invention. For example, a method includes

growing, in an appropriate medium, the cell, where the cell comprises a nucleic acid that comprises at least one selector codon and encodes a protein; and, providing the selected amino acid; where the cell further comprises: an orthogonal leucyl-tRNA (leucyl-O-tRNA) that functions in the cell and recognizes the selector codon; and, an orthogonal aminoacyl-

5   tRNA synthetase (O-RS) that preferentially aminoacylates the leucyl-O-tRNA with the selected amino acid. Typically, the leucyl-O-tRNA comprises at least about a 25%, 50%, 75%, 80%, 85%, 90%, 95% or 98% suppression activity in the presence of a cognate synthetase in response to a selector codon as compared to a control lacking the selector codon (and, typically, the cognate synthetase). A protein produced by this method is also a

10   feature of the invention.

[0158]   The invention also teaches variant orthogonal leucyl-tRNAs and variant orthogonal aminoacyl-tRNA synthetase species that display suppression activity, where the suppression activity is measured relative to the suppression activity of a leucyl-O-tRNA nucleotide sequence or an O-RS amino acid sequence provided by the present invention.

15   For example, the invention teaches variant leucyl-O-tRNA species that display suppression activity that is at least 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 98% or 99% as effective as a leucyl-O-tRNA sequence provided by the examples herein (e.g., SEQ ID NOs: 1-7 and 12). Similarly, the invention teaches variant O-RS species that display suppression activity that is at least 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 98% or

20   99% as effective as an O-RS sequence provided by the examples herein (e.g., SEQ ID NO: 15 and 16).

[0159]   In another aspect, the invention teaches variant orthogonal leucyl-tRNAs and variant orthogonal aminoacyl-tRNA synthetase species that display suppression activity that is equal to or greater than the suppression activity of a leucyl-O-tRNA nucleotide sequence or an O-RS amino acid sequence provided by the present specification. For example, the

25   invention teaches variant leucyl-O-tRNA species that display suppression activity that is at least 100% as effective as a leucyl-O-tRNA sequence provided by the examples herein (e.g., SEQ ID NO: 1-7 and 12).

[0160]   The compositions of the invention and compositions made by the methods of

30   the invention optionally are in a cell. The leucyl O-tRNA/O-RS pairs or individual components of the invention can then be used in a host system's translation machinery, which results in a selected amino acid, e.g., unnatural amino acid, being incorporated into a

protein. The International Application Number PCT/US2004/011786, filed April 16, 2004, and WO 2002/085923, entitled "IN VIVO INCORPORATION OF UNNATURAL AMINO ACIDS" describe this process, and is incorporated herein by reference. For example, when an leucyl O-tRNA/O-RS pair is introduced into a host, e.g., *Escherichia coli*, the pair leads to the in vivo incorporation of selected amino acid, such as an unnatural amino acid, e.g., a synthetic amino acid, such as derivative of a leucine amino acid, which can be exogenously added to the growth medium, into a protein, in response to a selector codon. Optionally, the compositions of the present invention can be in an in vitro translation system, or in an in vivo system(s).

[0161]   Essentially any protein (or portion thereof) that includes a selected amino acid, e.g., an unnatural amino acid, (and any corresponding coding nucleic acid, e.g., which includes one or more selector codons) can be produced using the compositions and methods herein. No attempt is made to identify the hundreds of thousands of known proteins, any of which can be modified to include one or more unnatural amino acid, e.g., by tailoring any available mutation methods to include one or more appropriate selector codon in a relevant translation system. Common sequence repositories for known proteins include GenBank EMBL, DDBJ and the NCBI. Other repositories can easily be identified by searching the internet.

[0162]   Typically, the proteins are, e.g., at least 60%, at least 70%, at least 75%, at least 80%, at least 90%, at least 95%, or at least 99% or more identical to any available protein (e.g., a therapeutic protein, a diagnostic protein, an industrial enzyme, or portion thereof, and the like), and they comprise one or more selected amino acid. Examples of therapeutic, diagnostic, and other proteins that can be modified to comprise one or more selected amino acid, e.g., an unnatural amino acid, can be found, but not limited to, those in International Application Number PCT/US2004/011786, filed April 16, 2004, and WO 2002/085923, entitled "IN VIVO INCORPORATION OF UNNATURAL AMINO ACIDS."

[0163]   In certain embodiments, the protein or polypeptide of interest (or portion thereof) in the methods and/or compositions of the invention is encoded by a nucleic acid. Typically, the nucleic acid comprises at least one selector codon, at least two selector codons, at least three selector codons, at least four selector codons, at least five selector

codons, at least six selector codons, at least seven selector codons, at least eight selector codons, at least nine selector codons, ten or more selector codons.

[0164]   Genes coding for proteins or polypeptides of interest can be mutagenized using methods well-known to one of skill in the art and described herein under "Mutagenesis and Other Molecular Biology Techniques" to include, e.g., one or more selector codon for the incorporation of a selected amino acid, e.g., an unnatural amino acid. For example, a nucleic acid for a protein of interest is mutagenized to include one or more selector codon, providing for the insertion of the one or more selected amino acids, e.g., unnatural amino acids. The invention includes any such variant, e.g., mutant, versions of any protein, e.g., including at least one selected amino acid. Similarly, the invention also includes corresponding nucleic acids, i.e., any nucleic acid with one or more selector codon that encodes one or more selected amino acid.

[0165]   To make a protein that includes a selected amino acid, one can use host cells and organisms that are adapted for the in vivo incorporation of the selected amino acid via orthogonal leucyl tRNA/RS pairs. Host cells are genetically engineered (e.g., transformed, transduced or transfected) with one or more vectors that express the orthogonal leucyl tRNA, the orthogonal leucyl tRNA synthetase, and a vector that encodes the protein to be derivatized. Each of these components can be on the same vector, or each can be on a separate vector, two components can be on one vector and the third component on a second vector. The vector can be, for example, in the form of a plasmid, a bacterium, a virus, a naked polynucleotide, or a conjugated polynucleotide.

Defining Polypeptides by Immunoreactivity

[0166]   Because the polypeptides of the invention provide a variety of new polypeptide sequences (e.g., comprising selected amino acids (e.g., unnatural amino acids) in the case of proteins synthesized in the translation systems herein, or, e.g., in the case of the novel synthetases, novel sequences of standard amino acids), the polypeptides also provide new structural features which can be recognized, e.g., in immunological assays. The generation of antisera, which specifically bind the polypeptides of the invention, as well as the polypeptides which are bound by such antisera, are a feature of the invention. The term "antibody," as used herein, includes, but is not limited to a polypeptide substantially encoded by an immunoglobulin gene or immunoglobulin genes, or fragments thereof which specifically bind and recognize an analyte (antigen). Examples include polyclonal,

-55-

monoclonal, chimeric, and single chain antibodies, and the like. Fragments of immunoglobulins, including Fab fragments and fragments produced by an expression library, including phage display, are also included in the term "antibody" as used herein. *See, e.g.,* Paul, *Fundamental Immunology*, 4th Ed., 1999, Raven Press, New York, for antibody structure and terminology.

[0167]   In order to produce antisera for use in an immunoassay, one or more of the immunogenic polypeptides is produced and purified as described herein. For example, recombinant protein can be produced in a recombinant cell. An inbred strain of mice (used in this assay because results are more reproducible due to the virtual genetic identity of the mice) is immunized with the immunogenic protein(s) in combination with a standard adjuvant, such as Freund's adjuvant, and a standard mouse immunization protocol (*see*, e.g., Harlow and Lane (1988) <u>Antibodies, A Laboratory Manual</u>, Cold Spring Harbor Publications, New York, for a standard description of antibody generation, immunoassay formats and conditions that can be used to determine specific immunoreactivity.

[0168]   Additional details on proteins, antibodies, antisera, etc. can be found in, e.g., International Application Number PCT/US2004/011786, filed April 16, 2004, WO 2002/085923, entitled "IN VIVO INCORPORATION OF UNNATURAL AMINO ACIDS;" International Application Numbers PCT/US2003/32870, filed October 15, 2003; and PCT/US2003/41346, filed December 22, 2003.

<u>KITS</u>

[0169]   Kits are also a feature of the invention. For example, a kit for producing a protein that comprises at least one selected amino acid, e.g., an unnatural amino acid, in a cell is provided, where the kit includes a container containing a polynucleotide sequence encoding an leucyl O-tRNA, and/or an leucyl O-tRNA, and/or a polynucleotide sequence encoding an leucyl O-RS, and/or an leucyl O-RS. In one embodiment, the kit further includes at least selected amino acid. In another embodiment, the kit further comprises instructional materials for producing the protein.

EXAMPLES

[0170]   The following examples are offered to illustrate, but not to limit the claimed invention. One of skill will recognize a variety of non-critical parameters that may be altered without departing from the scope of the claimed invention.

EXAMPLE 1: ADAPTATION OF AN ORTHOGONAL ARCHAEAL LEUCYL-TRNA AND SYNTHETASE PAIR FOR FOUR-BASE, AMBER, AND OPAL SUPPRESSION

[0171]   Recently, it has been shown that an amber suppressor tRNA-aminoacyl tRNA synthetase pair derived from the tyrosyl-tRNA synthetase of *Methanococcus jannaschii* can be used to genetically encode unnatural amino acids in response to the amber nonsense codon, TAG. This pair is unable to decode either the opal nonsense codon, TGA, or the four-base codon, AGGA. To overcome this, a leucyl-tRNA synthetase from *Methanobacterium thermoautotrophicum* and leucyl tRNA derived from *Halobacterium sp. NRC-1* was adapted as an orthogonal tRNA-synthetase pair in *E. coli* to decode amber (TAG), opal (TGA) and four-base (AGGA) codons. To improve the efficiency and selectivity of the suppressor tRNA, extensive mutagenesis was performed on the anticodon loop and acceptor stem. The two most significant criteria required for an efficient amber orthogonal suppressor tRNA are a CU(X)XXXAA anticodon loop and the lack of non-canonical or mismatched base pairs in the stem regions. These changes afford only weak suppression of TGA and AGGA. However, this information, together with an analysis of sequence similarity of multiple native archaeal tRNA sequences, led to efficient, orthogonal suppressors of opal codons and the four-base codon, AGGA. Ultimately, these additional orthogonal pairs can be used to genetically incorporate multiple unnatural amino acids into proteins.

[0172]   A great deal of effort has focused on the cotranslational incorporation of unnatural amino acids into proteins. Early work demonstrated that the translational machinery of *E. coli* would accommodate amino acids similar in structure to the common twenty (Hortin and Boime (1983) Methods Enzymol. 96, 777-784). This work was further extended by relaxing the specificity of endogenous *E. coli* synthetases so that they activate unnatural amino acids as well as their cognate natural amino acid. Moreover, it was shown that mutations in editing domains could also be used to extend the substrate scope of the endogenous synthetase (Doring et al., (2001) Science 292, 501-504). However, these strategies are limited to *recoding* the genetic code rather than *expanding* the genetic code and lead to varying degrees of substitution of one of the common twenty amino acids with an unnatural amino acid.

[0173] Later it was shown that unnatural amino acids could be site-specifically incorporated into proteins *in vitro* by the addition of chemically aminoacylated orthogonal amber

suppressor tRNAs to an *in vitro* transcription/translation reaction (Noren et al., (1989) <u>Science</u> 244, 182-188; Bain et al., (1989) <u>J. Am. Chem. Soc.</u> 111, 8013-8014; Dougherty (2000) <u>Curr. Opin. Chem. Biol.</u> 4, 645-652; Cornish et al., (1995) <u>Angew. Chem., Int. Ed.</u> 34, 621-633). It is clear from these studies that the ribosome and translation factors are

5   compatible with a large number of unnatural amino acids, even those with unusual structures. Unfortunately, the chemical aminoacylation of tRNAs is difficult, and this method can only produce microgram-scale quantities of protein due to the stoichiometric nature of the process. A catalytic *in vivo* method could overcome these limitations, and would also permit the study of proteins containing unnatural amino acids in living cells.

10   [0174] In order to add additional synthetic amino acids to the genetic code *in vivo* it is necessary to generate a 21st "orthogonal pair" of synthetase and tRNA that can function efficiently in the translational machinery. The synthetase should not cross-react with any of the endogenous tRNAs (40 in *E. coli*), and the orthogonal tRNA should not be aminoacylated by any of the endogenous synthetases (21 in *E. coli*). The tRNA should

15   decode only a specific new codon that is not decoded by any endogenous tRNA, and the synthetase should charge its tRNA with only a specific unnatural amino acid. We have successfully generated an orthogonal tRNA-synthetase pair from tyrosyl-tRNA synthetase of *Methanococcus jannaschii* which satisfies these requirements. This system has been used to incorporate a series of unnatural amino acids including keto amino acids (Wang et

20   al., (2003) <u>Proc. Natl. Acad. Sci. U. S. A.</u> 100, 56-61), photocrosslinking amino acids (Chin et al., (2002) <u>Proc. Natl. Acad. Sci. U. S. A.</u> 99, 11020-11024; Chin et al., (2002) <u>J. Am. Chem. Soc.</u> 124, 9026-9027), and heavy atom containing amino acids selectively into proteins in response to the TAG codon.

[0175] Several other orthogonal pairs have been reported. Glutaminyl (Liu and Schultz

25   (1999) <u>Proc. Natl. Acad. Sci. U. S. A.</u> 96, 4780-4785), aspartyl (Pastrnak et al., (2000) <u>Helv. Chim. Acta</u> 83, 2277-2286), and tyrosyl (Ohno et al., (1998) <u>J. Biochem. (Tokyo, Jpn.)</u> 124, 1065-1068; Kowal et al., (2001) <u>Proc. Natl. Acad. Sci. U. S. A.</u> 98, 2268-2273) systems derived from *S. cerevisiae* tRNAs and synthetases have been described for the potential incorporation of unnatural amino acids in *E. coli*. Systems derived from the *E. coli*

30   glutaminyl (Kowal et al., (2001) <u>Proc. Natl. Acad. Sci. U. S. A.</u> 98, 2268-2273) and tyrosyl (Edwards and Schimmel (1990) <u>Mol. Cell. Biol.</u> 10, 1633-1641) synthetase have been described for use in *S. cerevisiae*. The *E. coli* tyrosyl system can also function in mammalian cells and has been used for the incorporation of 3-iodo-L-tyrosine *in vivo*

(Sakamoto et al., (2002) <u>Nucleic Acids Res.</u> 30, 4692-4699). All of these systems have made exclusive use of the amber stop codon. To expand the genetic code beyond twenty-one amino acids, other orthogonal pairs and unique codons need to be identified.

[0176] A desired property of any orthogonal pair are a codon that is unique within the

5   genetic code and that will not cross-react with noncognate tRNAs. In addition to the amber stop codon (TAG), the opal nonsense codon (TGA) is one such candidate. A genetic code in which TAG and TGA encoded unnatural amino acids could encode 22 amino acids while preserving the ochre nonsense codon, UAA, which is the most abundant termination signal. The suppression of opal codons is robust *in vivo* but has not been frequently used for the

10   incorporation of unnatural amino acids *in vitro* due to high background readthrough of TGA codons (Cload et al., (1996) <u>Chem. Biol.</u> 3, 1033-1038). Another possible codon involves unnatural base pairs. Unnatural amino acids have been incorporated in response to novel codons containing the unnatural base (*iso*-dC)AG (Piccirilli et al., (1990) <u>Nature</u> 343, 33-37) or pyridin-2-one (Hirao et al., (2002) <u>Nat. Biotechnol.</u> 20, 177-182) using an *in vitro*

15   translation system. Adaptation of unnatural base pairs for the incorporation of unnatural amino acids into proteins *in vivo*, need the faithful replication and transcription of unnatural base pairs in DNA and RNA (Wu et al., (2002) <u>J. Am. Chem. Soc.</u> 124, 14626-14630). Another codon that can used to encode additional amino acids are four- and five-base codons. Using a library of tRNAs with randomized anticodon loops coupled with a

20   selection scheme, several highly efficient and non cross-reactive four- and five-base codons, were identified, including AGGA, UAGA, CCCU, and UAGA (Magliery et al., (2001) <u>J. Mol. Biol.</u> 307, 755-769; Anderson et al., (2002) Chem. Biol. 9, 237-244).

[0177] Regardless of the codon chosen, it is useful to generate additional orthogonal tRNA-synthetase pairs that can translate these codons with high fidelity and good efficiency.

25   Because the tRNA anticodon loop is a major identity element for recognition by most synthetases, one must identify a synthetase that does not recognize these identity elements in order to generate suppressor tRNAs for these unusual codons. The leucyl-, seryl-, and alanyl-tRNA synthetases of *E. coli* are well known to tolerate extensive substitutions in the anticodon loop (Shimizu et al., (1992) <u>J. Mol. Evol.</u> 35, 436-443; Kleina (1990) <u>J. Mol.</u>

30   <u>Biol.</u> 213, 705-717; Sampson and Saks (1993) <u>Nucleic Acids Res.</u> 21, 4467-4475). Some homologous archaeal or eukaryotic synthetases may have similar properties. Herein are derivatives of a leucyl-tRNA synthetase from *Methanobacterium thermoautotrophicum* and leucyl tRNAs derived from *Halobacterium sp. NRC-1* that act as orthogonal tRNA-

synthetase pairs for the amber codon in *E. coli*. Moreover, information gained in these studies, together with multiple sequence alignments of native archaeal tRNA sequences, allowed us to design efficient orthogonal suppressor tRNAs of opal codons and a four-base codon, AGGA.

Material and Methods

[0178] *Strains, plasmids, and materials.* All *in vivo* manipulations were carried out in *E. coli* strain TOP10 (Invitrogen) in LB media at 37°C. *Halobacterium sp. NRC-1* was purchased from the American Type Culture Collection (ATCC). PCR was carried out according to standard protocols with a mixture of *Taq* (Promega) and *Pfu* (Stratagene) polymerases. Oligonucleotides were synthesized by Genosys, Operon, or the UCSF Biomolecular Resource Center. For oligonucleotides containing degenerate bases, the phosphoramidites were premixed to avoid bias. Standard protocols were employed for subcloning with restriction enzymes (NEB) and T4 DNA ligase (NEB). Plasmids were introduced into *E. coli* by electroporation. Sequence analysis was performed using the Genetics Computer Group, Inc. (GCG) software. The sequences of all plasmids were confirmed by restriction mapping and sequencing.

[0179] *Cloning of tRNA synthetase genes.* Genomic DNA was either purchased from ATCC or was prepared from a cell pellet purchased from ATCC. Genomic DNA was extracted using the DNeasy kit (Qiagen). Synthetase genes were amplified from genomic DNA by PCR then subcloned into the *Nco*I and either *Eco*RI, *Kpn*I, or *Pvu*II sites of plasmid pKQ. More details on the cloning of these genes can be found Table 2 and is also available on the Internet at http://pubs.acs.org. Plasmid pKQ contains the ribosome binding site, multiple cloning site, and *rrnB* terminator from plasmid pBAD-Myc/HisA (Invitrogen) under control of a constitutive glutamine promoter. The plasmid also contains a ColE1 origin of replication, and a kanamycin resistance gene for plasmid maintenance.

[0180] *Constructions of reporter plasmids.* Beta-lactamase reporter plasmids were constructed from plasmid pACKO-Bla. This plasmid was constructed with a p15a origin, a chloramphenicol resistance gene, and unique sites for insertion of a gene for β-lactamase and a tRNA under control of the strong, constitutive *lpp* promoter. Site A184 of the β-lactamase gene was changed to TAG, AGGA, or TGA by an overlap PCR strategy, and the genes were subcloned into the *Aat*II and *Xma*I sites of pACKO-Bla to give plasmids pACKO-A184TAG, pACKO-A184AGGA, and pACKO-A184TGA.

[0181]   *Constructions of tRNA plasmids.* Genes for individual tRNAs and for tRNA libraries were constructed by extension reactions and subcloned into the *EcoRI* and *PstI* sites of pACKO-Bla derivatives. All libraries represented at least 10-fold more members than the theoretical size of the library to ensure high coverage.

5   [0182]   *Measurement of suppression efficiency.* A series of LB agar plates were prepared with 25 µg/mL of kanamycin, 25 µg/mL of chloramphenicol, and concentrations of ampicillin between 5 and 1000 µg/mL. Synthetase and tRNA plasmids were cotransformed and plated at densities below 100 cells per plate. Suppression efficiency was reported as the highest concentration at which cells survived to form colonies among a

10   series of plates for which the next highest and lowest concentrations would be within 20% of the reported value.

[0183]   *Selection of libraries and characterization of selectants.* All tRNA libraries were subjected to ampicillin selection and the surviving colonies were isolated and sequenced by the method described previously (Magliery et al., (2001) J. Mol. Biol. 307,

15   755-769). Briefly, libraries were spread on LB plates containing 25 µg/mL of kanamycin and chloramphenicol for plasmid maintenance, and varying concentrations of ampicillin for selection. After 24 hours of growth, the plates were scraped, and the cells were diluted slightly then spread again on ampicillin plates. After colonies appeared, plates were again scraped and plated at dilute cell densities on a range of plates with different ampicillin

20   concentrations. Selectants were isolated, sequenced, and then confirmed by retransformation into cells containing synthetase-expressing plasmids.

[0184]   *Beta-galactosidase reporter assays.* The full-length *lacZ* gene of plasmid pBAD-Myc/His/lacZ (Invitrogen) was amplified by PCR and subcloned into plasmid pLASC to obtain plasmid pLASC-lacZ. This pSC101-derived plasmid expresses *lacZ* gene

25   under the control of an *lpp* promoter and has an ampicillin resistance gene for plasmid maintenance. Derivatives of this plasmid were constructed wherein Leu-25 of the peptide VVLQRRDWEN of *lacZ* was replaced by TAG, TGA, or AGGA codons, or sense codons for tyrosine, serine, or leucine. The appropriate pLASC-lacZ-, pACKO-Bla-, and pKQ-derived plasmids were cotransformed and grown to an OD_{600} of 0.5. Beta-galactosidase

30   assays were performed in quadruplicate using the BetaFluor™ β-Galactosidase Assay Kit (Novagen). Percent suppression was calculated as the percentage of activity for a sample

relative to the value observed from the pLASC-lacZ construct with the corresponding sense
codon at position 25. Cells containing pLASC-lacZ plasmids with sense codons at position
25 were also assayed by 2-nitrophenyl-β-D-galactopyranoside assays (Miller (1972)
Experiments in molecular genetics, Cold Spring Harbor Laboratory, Cold Spring Harbor,
N.Y.), and activity was calculated in Miller units.

[0185]   *Purification of synthetase proteins.* Synthetase genes were cloned in frame
with the C-terminal myc/his tag of pBAD-Myc/HisA (Invitrogen). Protein purification was
performed with the Qiaexpressionist kit (Qiagen) by the manufacturer's protocol under
native conditions. Protein concentrations were measured by the BCA Protein Assay Kit
(Pierce) and analyzed by SDS-PAGE.

[0186]   *In vitro aminoacylation assays.* Aminoacylation assays were performed by
methods described previously (Hoben and Soll (1985) Methods Enzymol. 113, 55-59) in
20 µL reactions containing 50 mM Tris-Cl, pH 7.5, 30 mM KCl, 20 mM $MgCl_2$, 3 mM
glutathione, 0.1 mg/mL BSA, 10 mM ATP, 1 µM (79 Ci/mmol) [^{3}H] leucine (Amersham),
750 nM synthetase, and 0, 2, 10, or 40 µM crude total tRNA. Crude total *E. coli* tRNA was
purchased from Roche, and halobacterial tRNA was extracted from cultures of
Halobacterium sp. NRC-1 with the RNA/DNA Extraction Kit (Qiagen).

[0187]   Detailed information for the cloning of archaeal leucyl-tRNA synthetases can
be found below and is also available on the Internet at http://pubs.acs.org.

TABLE 2: CLONING OF ARCHAEAL LEUCYL-tRNA SYNTHETASES INTO PLASMID pKQ

Organism	Accession number	ATCC Number	Forward Oligo	Reverse Oligo
Halobacterium sp. NRC-1	NP_280869.1	700922	ca214	ca215
Escherichia coli (strain HB101)	P07813	N/A	ca244	ca215
Methanococcus jannaschii	Q58050	43067D	ca246	ca247
Archaeoglobus fulgidus	O30250	49558D	ca261	ca247
Aeropyrum pernix K1	Q9YD97	700793D	ca263	ca264
Pyrococcus horikoshii	O58698	700860	ca265	ca266
Methanobacterium thermoautotrophicum	O27552	700791	ca274	ca275

Organism	Restriction enzymes
Halobacterium sp. NRC-1	NcoI/EcoRI
Escherichia coli (strain HB101)	NcoI/EcoRI
Methanococcus jannaschii	NcoI/KpnI
Archaeoglobus fulgidus	NcoI/KpnI

Aeropyrum pernix K1	BspHI/EcoRI (subcloned into NcoI/EcoRI sites)
Pyrococcus horikoshii	NcoI/EcoRV (subcloned into NcoI/PvuII sites)
Methanobacterium thermoautotrophicum	BsmBI (subcloned into NcoI/EcoRI sites)

Oligo Sequences

ca214	GGTTTCCATGGGAGAGCAAGCCACCTAC
ca215	GGTTTGGAATTCAGTCGTCGGCTTCGTCG
ca244	CGAAACCATGGAAGAGCAATACCGCCCGGAAG
ca245	CCAAAGAATTCCCGCCAACGACCAGATTGAGGAG
ca246	CGAAACCATGGTTATGATTGACTTTAAAG
ca247	CGAAAGGTACCTTGTATTCAAGATAAATAGCTGG
ca261	GCGAACCATGGGCGATTTCAGGATAATTGAG
ca262	CAATTGGTACCTTAAGCAACATAAATCGCG
ca263	GGATTATCATGAAGCGACTAAAGGCCGTGGAGGAG
ca264	CACTTGAATTCTTAGCCTCCTCTCTTCTCCGC
ca265	CGAATCCATGGCTGAGCTTAACTTCAAGG
ca266	GGATGGATATCACTCGATGAAGATGGCAG
ca274	GGAGACGTCTCTCATGGATATTGAAAGAAAATGGCG
ca275	CGTTACGTCTCGAATTGGAAAAGAGCTGTCTGAGG

Results and Discussion

5　**[0188]**　　　　*Identification of orthogonal tRNAs.* Previous studies (Kwok and Wong (1980) Can. J. Biochem. 58, 213-218) have shown that halobacterial tRNAs are inefficiently charged by the *E. coli* leucyl-tRNA synthetase. The similarity between the halobacterial and other archaeal leucyl-tRNAs (see **Figure 1, Panel C**) led us to believe that tRNAs from other archaeans can also be orthogonal to the *E. coli* synthetases. The sequences were

10　chosen to broadly represent the family of archaeal leucyl-tRNAs and included $tRNA_3^{Leu}$ of *Archaeoglobus fulgidus* (AfL3), $tRNA_4^{Leu}$ of *Halobacterium sp. NRC-1* (HhL4), $tRNA_2^{Leu}$ of *Methanococcus jannaschii* (MjL2), $tRNA_5^{Leu}$ of *Pyrococcus furiosus* (PfL5), and $tRNA_2^{Leu}$ of *Pyrococcus horikoshi* (PhL2) (see **Figure 1, Panel C** for sequences). In all cases, the anticodon was changed to CUA, and CCA was added to the 3' terminus if the

15　sequence was not present in the source gene to obtain an amber suppressor tRNA.

[0189]　　　　To measure the activity of suppressor tRNAs, a selection system was developed based on the *in vivo* suppression of nonsense or frameshift mutations introduced into the gene for β-lactamase *(bla)*. Reporter genes for *bla* variants with TAG, AGGA, and TGA at position A184 (a permissive site (Liu and Schultz (1999) Proc. Natl. Acad. Sci. U.

20　S. A. 96, 4780-4785)) were constructed in plasmid pACKO-Bla, a medium copy plasmid

derived from pACYC184. Bacteria transformed with these reporter constructs are unable to grow on LB agar plates with ampicillin concentrations greater than 5 μg/mL, only slightly higher than the value (2 μg/mL ampicillin) observed for bacteria transformed with no plasmids. Plasmids derived from pACKO-Bla can also express tRNA genes under the

5   control of a strong *lpp* promoter. When the robust amber suppressor gene *supD*, a tRNA efficiently charged by *E. coli* seryl tRNA synthetase, is expressed from pACKO-A184TAG (which encodes the A184TAG variant of *bla*), host bacteria survive at an ampicillin concentration of 1000 μg/mL. In contrast, in the case of an orthogonal tRNA, which cannot be efficiently charged by endogenous *E. coli* synthetases, ampicillin resistance should be

10   less than 5 ug/mL. Conversely, if the tRNA is not orthogonal, or if a heterologous synthetase capable of charging the tRNA is co-expressed in the system, a higher level of ampicillin resistance should be observed.

[0190]   The genes for the five potential orthogonal amber suppressor tRNAs were integrated into pACKO-A184TAG. *E. coli* hosts expressing the HhL4-derived suppressor,

15   designated HL(TAG)1, could survive to only 5 μg/mL ampicillin, the MjL2- and PhL2-derived suppressors to 7 μg/mL, and the PfL5- and AfL3-derived suppressors to 20 μg/mL ampicillin. Therefore, all five suppressor tRNAs are either weak suppressor tRNAs or are inefficiently charged by *E. coli* aminoacyl-tRNA synthetases.

[0191]   *Cloning of archaeal leucyl-tRNA synthetases.* Due to the high homology of

20   the archaeal leucyl-tRNAs, we anticipated that the archeal leucyl-tRNA synthetases might have similar tRNA recognition properties. Therefore both species specific and cross species combinations of archeal leucyl-tRNAs and synthetases were examined in order to find an optimal pair for use in *E. coli*. The leucyl-tRNA synthetases from *Archaeoglobus fulgidus* (AfLRS), *Aeuropyrum pernix* (ApLRS), *Halobacterium sp. NRC-1* (HhLRS),

25   *Methanococcus jannaschii* (MjLRS), *Methanobacterium thermoautotrophicum* (MtLRS), and *Pyrococcus horikoshi* (PhLRS) were chosen as initial candidates due to the availability of the genome sequences and commercial availability of the organisms. The genes for these synthetases were cloned under the control of a constitutive glutamine promoter on the high copy plasmid, pKQ, which was constructed from pBR322 and contains a kanamycin

30   resistance gene. The *leuS* gene from *E. coli* (EcLRS) was also cloned as a negative control. Synthetase expression plasmids and reporter constructs were cotransformed and assayed for activity by ampicillin selection (**Figure 2**). In general, the reporter plasmid containing the

HhL4 suppressor tRNA, HL(TAG)1, gave the largest enhancement in suppression efficiency upon cotransformation with synthetase-expressing plasmids, but the PhL2- and AfL3-derived tRNAs also show a suppression enhancement. The MjL2- and PfL5-derived suppressor tRNAs survive to the same concentrations of ampicillin regardless of whether or

5 not the archaeal synthetase is present, and were not pursued further. From all 35 combinations of synthetase and reporter plasmids, the highest levels of ampicillin resistance result when the synthetases, MtLRS or MjLRS, are expressed with the HhL4-derived suppressor tRNA. The AfLRS construct gives slightly lower levels of resistance, and all other synthetases give no increase in suppression efficiency over background levels. With

10 MjLRS or MtLRS, cells expressing HL(TAG)1 survive to 35 µg/mL ampicillin, but only 5 µg/mL with the *E. coli* synthetase or plasmid lacking the synthetase. Cells expressing AfLRS can survive to 25 µg/mL ampicillin when coexpressed with HL(TAG)1. From these *in vivo* suppression screens, three synthetases, MtLRS, MjLRS, and AfLRS, were identified as candidates for an orthogonal pair with the HhL4-derived amber suppressor tRNA,

15 HL(TAG)1.

[0192]   *In vitro charging assays.*   An *in vivo* suppression screen can distinguish active and inactive aminoacyl-tRNA synthetases, but it cannot distinguish an orthogonal synthetase from one that cross-reacts with *E. coli* tRNA. To determine the permissiveness of AfLRS, MjLRS, and MtLRS for *E. coli* tRNA, the synthetases were overexpressed,

20 purified, and then subjected to *in vitro* aminoacylation assays to measure their ability to charge *E. coli* tRNA. AfLRS, MjLRS, and MtLRS were purified from an arabinose promoter over-expression system by Ni-NTA affinity chromatography in yields of 14, 8, and 3 mg/L respectively. *In vitro* aminoacylation assays were performed with tritium-labeled leucine and either *E. coli* or *Halobacterium NRC-1* total tRNA (**Figure 3, Panels A**

25 **and B**). Based on the charging of 10 µM crude total tRNA, MtLRS and AfLRS charge halobacterial tRNA 54- and 21- fold more efficiently than *E. coli* tRNA, respectively. The MjLRS enzyme, however, shows only a 6-fold preference for halobacterial tRNA. The *E. coli* enzyme was 100-fold more efficient at charging *E. coli* crude total tRNA than halobacterial tRNA. Therefore, MtLRS and AfLRS are good candidates for orthogonal

30 aminoacyl-tRNA synthetases with respect to *E. coli* tRNA, but MjLRS is not. Since MtLRS showed a higher level of suppression with HL(TAG)1 *in vivo* than did AfLRS, the

MtLRS/HL(TAG)1 pair was carried forward as a potential new orthogonal pair for use in *E. coli.*

[0193]   *Optimization of the tRNA anticodon loop.* The robust endogenous amber suppressor *supD* confers survival to 1000 µg/mL ampicillin when expressed from pACKO-

5   A184TAG. In contrast, cells expressing the MtLRS/HL(TAG)1 pair survive to only 35 µg/mL ampicillin, which corresponds to a 2.9% suppression efficiency as determined from β-galactosidase assays (Table 1). We therefore sought to improve the activity of the system. Previous experiments on frameshift, missense, and nonsense suppression revealed that A37 was a highly conserved feature in robust suppressor tRNAs (Magliery et al.,

10   (2001) J. Mol. Biol. 307, 755-769). HhL4 has a G at position 37, therefore substitution of G37 to A might be expected to improve suppression efficiency. To examine this and other possible anticodon loop mutants, a library was constructed in which the 7 positions of the anticodon loop (positions 32-38, see **Figure 4, Panel A**) in HhL4 were replaced with degenerate bases and subcloned into pACKO-A184TAG. The library of tRNAs was

15   cotransformed with pKQ-MtLRS and subjected to ampicillin selection initially at 35 µg/mL ampicillin for two rounds of selection, then plated on a series of plates with increasing ampicillin concentration in the third round of selection. At the highest concentration of ampicillin for which growth was observed (500 µg/mL), the only clone found had an anticodon loop with the sequence CUCUAAA, corresponding to a simple G37A mutation

20   (Table 1). When cotransformed with pKQ-MtLRS, this clone could survive to 500 µg/mL ampicillin. In the absence of the synthetase it survived to only 25 µg/mL ampicillin. Under similar conditions, cells containing the wild-type *M. jannaschii* tyrosyl orthogonal amber suppressor tRNA survive to 350 µg/mL ampicillin in the presence of the cognate synthetase and to 60 ug/mL ampicillin without the synthetase.

Table 1. Suppression efficiency of mutant orthogonal tRNAs.

Reporter Plasmid	Miller Units		
pLASC-lacZ(Leu)	210	±	2
pLASC-lacZ(Ser)	200	±	5
pLASC-lacZ(Tyr)	192	±	7
pLASC-lacZ(TAG)	1	±	1
pLASC-lacZ(AGGA)	2	±	1
pLASC-lacZ(TGA)	1	±	1

	Percent Suppression[a]						
Suppressor tRNA	with pKQ			with synthetase			Sequence
HL(TAG)1	0.4	±	0.1%	2.9	±	0.8%	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACTCTAGATCCGTTCTCGTAGGA GTTCGAGGGTTCGAATCCCTTCCCTCGCACCA
HL(TAG)2	0.3	±	0.1%	9.6	±	0.4%	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACTCTAAATCCGTTCTCGTAGGA GTTCGAGGGTTCGAATCCCTTCCCTCGCACCA
HL(TAG)3	1.5	±	1.2%	33.2	±	4.4%	CCCAGGGTAGCCAAGCTCGGCCAACGGCGACGGACTCTAAATCCGTTCTCGTAGGA GTTCGAGGGTTCGAATCCCTTCCCTGGGACCA
HL(AGGA)1	0.4	±	0.1%	4.6	±	2.1%	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACTTCCTAATCCGTTCTCGTAGG AGTTCGAGGGTTCGAATCCCTTCCCTCGCACCA
HL(AGGA)2	0.7	±	0.3%	14.9	±	6.1%	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACTTCCTAATCCGTTCTCGTAGG AGTTCGAGGGTTCGAATCCCTCCCCTCGCACCA
HL(AGGA)3	7.4	±	0.4%	35.5	±	1.4%	GCGGGGGTTGCCGAGCCTGGCCAAAGGCGCCGGACTTCCTAATCCGGTCCCGTAGG GGTTCCGGGGTTCAAATCCCCGCCCCCGCACCA
HL(TGA)1	4.7	±	1.5%	60.8	±	7.0%	GCGGGGGTTGCCGAGCCTGGCCAAAGGCGCCGGACTTCAAATCCGGTCCCGTAGGG GTTCCGGGGTTCAAATCCCCGCCCCCGCACCA
J17[b]	0.2	±	0.1%	18.5	±	4.8%	CCGGCGGTAGTTCAGCAGGGCAGAACGGCGGACTCTAAATCCGCATGGCGCTGGTT CAAATCCGGCCCGCCGGACCA
SupD	42.8	±	7.1%	ND			GGAGAGATGCCGGAGCGGCTGAACGGACCGGTCTCTAAAACCGGAGTAGGGGCAAC TCTACCGGGGGTTCAAATCCCCCTCTCTCCGCCA
Ser2AGGA	25.2	±	0.1%	ND			GGAGAGATGCCGGAGCGGCTGAACGGACCGGTCTTCCTAAACCGGAGTAGGGGCAA CTCTACCGGGGGTTCAAATCCCCCTCTCTCCGCCA

5

[a]β-Galactosidase activity was determined for tRNA reporter plasmids derived from pACKO-Bla cotransformed with the appropriate pLASC-lacZ mutant and either a synthetase-expressing plasmid or a plasmid with no synthetase. Activity is reported as the percentage of activity observed relative to the value observed from the pLASC-lacZ construct with a leucyl (wild-type), seryl, or tyrosyl sense codon at position 10   25. In each case, the codon at position 25 of *lacZ* is designated in parentheses.

[b]J17, the *M. jannaschii* tyrosyl amber suppressor tRNA with improved orthogonality (Wang and Schultz (2001) Chem. Biol. 8, 883-890) was expressed in plasmid pACKO-A184TAG in the presence of pLASC-lacZ(TAG) and either pKQ or pBK-JYRS.

15   **[0194]**   *Randomization of leucyl acceptor stem.* Although the activity of the HhL4-derived amber tRNA was significantly improved with the G37A mutation, the suppression level in the absence of the synthetase increased from 5 to 25 μg/mL ampicillin. To overcome the undesired increase in background suppression, a mutant of the HhL4-derived tRNA was sought that would not cross react with *E. coli* aminoacyl-tRNA synthetases.

20   Almost all of *E. coli* synthetases recognize bases within the acceptor stem of their cognate tRNAs. Therefore, we anticipated that mutations within this region of the tRNA might eliminate interactions between the orthogonal tRNA and the cross-reactive synthetase. A

library in which the 3 terminal base pairs of the acceptor stem and the discriminator base were randomized (positions 1-3 and 70-73, the randomized region is outlined in **Figure 4, Panel A**) was constructed from the HL(TAG)2 mutant tRNA and subcloned into pACKO-A184TAG.

5   **[0195]**   To identify members of this tRNA library that retained activity but were even poorer substrates for endogenous synthetases, a selection strategy was adopted from previous work on the *M. jannaschii* system (Wang and Schultz (2001) <u>Chem. Biol.</u> 8, 883-890). To isolate a pool of mutant tRNAs that had comparable activity to the G37A mutant of the HhL4-derived tRNA, the tRNA library in which the acceptor stem was randomized

10   was cotransformed with pKQ-MtLRS and subjected to two rounds of positive selection at 500 µg/mL ampicillin. Six clones surviving the positive selection were sequenced, and all were unique and conserved the discriminator base, A73 (**Figure 4, Panels A and B**). In all cases the stem positions had standard Watson-Crick base pairs. To identify members of the pool of active clones that would not be charged by endogenous aminoacyl-tRNA

15   synthetases, the surviving tRNA-expressing plasmids were transferred into cells containing a barnase reporter plasmid, pSCB2. This plasmid contains the gene for the RNase, barnase, with two TAG codons at permissive positions 2 and 44, under control of the arabinose promoter, as well as the gene for β-lactamase. Any tRNA that is aminoacylated by an endogenous *E. coli* synthetase will result in suppression of the nonsense codons and cell

20   death. The cells were plated on LB plates containing 25 µg/mL of chloramphenicol, 50 µg/mL of ampicillin to maintain the plasmids, and 0.2% arabinose to induce expression of the barnase gene. Sixteen survivors were sequenced, and three unique sequences were identified. All three clones had reversed the 3:70 base pair from G:C to C:G. Of these, mutant HL(TAG)3 gave the highest level of suppression in the presence of MtLRS

25   (600 µg/mL ampicillin) and only survived to 7.5 µg/mL ampicillin without the synthetase. These values correspond to 33.2% suppression in the presence of MtLRS and 1.5% in the absence of the synthetase as determined by β-galactosidase amber suppression assays (see Table 1). By comparison, the mutant *M. jannaschii* suppressor tRNA, J17, gives values of 18.5% and 0.2% with and without the *M. jannaschii* tyrosine synthetase, respectively.

30   **[0196]**   *Identification of AGGA suppressors.* To expand the list of codons that can be used for unnatural amino acid mutagenesis, a tRNA that could efficiently suppress a four-base codon was sought. Previous studies indicated that the four-base codon AGGA

can be efficiently suppressed in *E. coli*, and tRNAs with 8 nucleotide anticodon loops were
the most efficient suppressors of AGGA codons (Magliery et al., (2001) <u>J. Mol. Biol.</u> 307,
755-769). A β-lactamase reporter plasmid analogous to the TAG reporter was constructed
but with A184 replaced by AGGA instead of TAG. Normal translation in the absence of a

5   +1 frameshift suppressor tRNA should result in missense errors downstream of position 184
and premature truncation of the protein. A library of tRNAs derived from the HhL4 tRNA
was constructed in which the 7 nucleotide anticodon loop was replaced with 8 random
nucleotides. The library was subcloned into pACKO-A184AGGA, cotransformed with
pKQ-MtLRS, and then subjected to ampicillin selection. At the highest concentration of

10   ampicillin at which growth was observed, 75 μg/mL, only one clone, HL(AGGA)1, was
found. This clone had the anticodon loop sequence CUUCCUAA. As was the case with
the *bla* A184TAG reporter plasmid, cells transformed with pACKO-A184AGGA can
survive to only 5 μg/mL ampicillin in the absence of a suppressor tRNA. Therefore, the
clone identified, HL(AGGA)1, is a weak suppressor of AGGA codons.

15   [0197]   During these experiments, serendipitous mutants capable of surviving up to
300 μg/mL ampicillin were identified. These mutants were no longer orthogonal, and all
had multiple point mutations relative to the parent sequence. All of the clones contained the
substitution T65C. This mutation corrects the G:U mismatch present in the TψC loop stem,
suggesting that this G:U base pair might be detrimental to suppressor activity. We therefore

20   decided to randomize this base; a library was made in which the 49:65 base pair was
randomized in HL(AGGA)1. The library was subcloned into pACKO-A184AGGA, and
then cotransformed with pKQ-MtLRS. Of the 16 library members, the most efficient
suppressor, HL(AGGA)2, was identified by ampicillin selection. This clone contained a
T65C mutation and could survive to 125 μg/mL ampicillin. Nevertheless, this level of

25   activity was far lower than that observed for the corresponding amber suppressors.
Consequently, alternative strategies were considered.

[0198]   Mutations in the D-loop have been previously implicated in frameshift
suppression (Tuohy et al., (1992) <u>J. Mol. Biol.</u> 228, 1042-1054), and we next hypothesized
that such mutations might improve the suppression efficiency of the AGGA suppressor.

30   Libraries wherein the 13 nucleotides of the D-loop (position 14-21, see **Figure 4, Panel A**)
were replaced with 11 or 13 random nucleotides were prepared in pACKO-A184AGGA.
Although a great deal of sequence diversity was observed among the survivors at the

highest concentrations of ampicillin (125 μg/mL), no mutants were observed with increased activity relative to the parent tRNA.

[0199]   *A consensus-derived AGGA suppressor tRNA.* In examining the sequence of the HhL4-derived tRNA, there was no obvious explanation for the poor activity of this

5   suppressor. Rather than mutate HL(AGGA)2 further, we pursued an alternative approach. The archaeal leucyl-tRNAs are highly similar, varying from each other usually by only a few base substitutions (**Figure 1, Panel C**). The entire family would be well represented by a library derived from a consensus sequence with many random mutations throughout. The consensus sequence was compiled with the GCG program *pileup*, and those positions

10   considered degenerate by the program were changed to the most frequent base at those positions. The anticodon loop was changed to CUUCCUAA since this sequence was already shown to be the optimal sequence for an AGGA suppressor derived from HhL4. The final sequence used as the consensus sequence is shown in **Figure 5**. A library was synthesized by overlap extension of oligonucleotides in which each site of the tRNA gene

15   was synthesized as a doped mixture of 90% the consensus sequence and 10% a mixture of the other 3 bases. The library was subcloned into pACKO-A184AGGA. Sequencing of 24 naïve clones revealed that the average number of mutations per clone was 5.9, and these mutations were randomly distributed throughout the tRNA sequence. After cotransformation with pKQ-MtLRS and selection on ampicillin plates, several clones

20   survived to 300 μg/mL of ampicillin and were found to be the original sequence with the 27:42 and 49:65 base pairs changed to the canonical base pairs T27:A42, G27:C42, or C27:G42, and G47:C65 or C47:G65 (**Figure 5**). The most efficient suppressor, designated HL(AGGA)3, can survive to 300 μg/mL ampicillin in the presence of pKQ-MtLRS but to only 30 μg/mL in the absence of the synthetase, which correspond to 35.5% and 7.4%

25   suppression, respectively, as determined by β-galactosidase assays (Table 1).

[0200]   *Identification of opal suppressor tRNAs.* To further expand the list of codons, we sought opal suppressors derived from HhL4. A reporter plasmid, pACKO-A184TGA, was constructed in which the A184 position of β-lactamase was changed to TGA. This *bla* A184TGA reporter plasmid can survive to 10 μg/mL ampicillin without any

30   suppressor tRNA present, whereas the TAG and AGGA reporters could survive to only 5 μg/mL. In the case of opal suppression, there is background read-through that leads to the

production of a small amount of protein even in the absence of a suppression system. Nevertheless, this level is quite small. To identify suppressors, a library in which the anticodon loop (positions 31-38) of HhL4 was replaced with 7 degenerate nucleotides was prepared in pACKO-A184TGA. When cotransformed with pKQ-MtLRS, no members of

5   this library could survive on ampicillin plates at 50 μg/mL. Instead of HhL4, a library was prepared in which the 8 nucleotide anticodon loop was randomized with 7 nucleotides in HL(AGGA)3, the most robust AGGA suppressor identified from the consensus sequence. At the highest concentrations of ampicillin at which growth was observed (300 μg/mL) only one clone, designated HL(TGA)1, with the sequence CUUCAAA was found. The clone

10   can survive to 350 μg/mL ampicillin when coexpressed with pKQ-MtLRS, but can survive to only 30 μg/mL without the synthetase plasmid, which corresponds to 60.8% suppression as determined by β-galactosidase assays (Table 1). Apparently, the beneficial effects of using the consensus sequence are not limited to frameshift suppression.

[0201]   *Identification of new orthogonal pairs.* One approach to construct

15   orthogonal tRNA-synthetase pairs is to adapt eukaryotic or archaeal synthetases and tRNAs for use in *E. coli.* Several yeast synthetases, notably glutamine, aspartic acid, arginine, and tyrosine, have been shown not to recognize *E. coli* tRNA, and might therefore be useful for the construction of orthogonal tRNA-synthetase pairs.. Unfortunately, many eukaryotic synthetases express poorly or have low specific activity in *E. coli.* Eukaryotic synthetases,

20   particularly the mammalian enzymes, are often organized into large complexes (Mirande et al., (1982) EMBO J. 1, 733-736), and the low activity often observed may be related to the inability to form these complexes in *E. coli.*

[0202] The success of the *M. jannaschii* tyrosyl orthogonal pair (Wang et al., (2001) Science 292, 498-500) suggested that archaebacteria may in general be a good source of

25   orthogonal pairs. Early work on the halophile *Halobacterium cutirebrum* (Kwok and Wong (1980) Can. J. Biochem. 58, 213-218) indicated that almost all the tRNAs of this archaean (notably leucine, arginine, tyrosine, serine, histidine, and proline) cannot be charged by *E. coli* aminoacyl-tRNA synthetases. Indeed, archaeal tRNA synthetases are more similar to their eukaryotic than prokaryotic counterparts in terms of homology and tRNA recognition

30   elements. Unlike their eukaryotic counterparts, however, there is currently no evidence for their higher order assembly into structured multimers (Tumbula et al., (1999) Genetics 152, 1269-1276; Woese et al., (2000) Microbiol. Mol. Biol. Rev. 54, 202-236). Moreover, since

most archaea are thermophiles, active synthetases from archaea can be expressed in good yields in *E. coli* and can be readily purified in active form. Due to extensive sequencing efforts, at least 16 archaeal genome sequences are currently available, which together with the lack of introns in the genome, greatly facilitates the PCR amplification of the archaeal

5   synthetase genes. For all of the above reasons, our attention has focused on the archaea as a source for orthogonal pairs.

[0203]   Another design issue in the construction of orthogonal tRNA-synthetase pairs is the ability of the aminoacyl-tRNA synthetase to recognize mutants of the cognate tRNA with altered anticodon loops (i.e., nonsense or missense suppressors). Aminoacyl-

10   tRNA synthetases frequently use the anticodon loop as a major positive identity element, and mutations in this region of the tRNA frequently result in impaired synthetase recognition. The leucyl-tRNA synthetases frequently lack strong anticodon recognition elements, and a leucyl orthogonal tRNA-synthetase pair can therefore be able to decode a variety of codons, including amber, opal and four-base codons. Of the archaeal leucyl-

15   tRNA synthetases, only the enzyme from *Haloferax volcanii* has been thoroughly investigated (Soma et al., (1999) J. Mol. Biol. 293, 1029-1038). The synthetase does not recognize bases in the anticodon loop; instead, a highly conserved pattern of mismatches within the variable loop is the primary recognition element for the synthetase. Although the cloning of the gene for this enzyme has not been reported, the sequenced genome of a

20   closely related archaean, *Halobacterium sp. NRC-1*, is available. A multiple sequence alignment of leucyl-tRNA synthetases from many phyla including archaeal, prokaryotic, and eukaryotic sequences (**Figure 1, Panel A**) shows that the halophilic enzyme is unusual among the family of archaeal synthetases, having greater homology to the prokaryotic branch than the eukaryotic or archaeal branches. Unlike the synthetases, all archaeal leucyl

25   tRNAs are highly homologous and share absolutely conserved features such as A73, G37, and a 12 nucleotide variable loop with 2 unpaired bases (**Figure 1, Panel B**). The conservation of these positive recognition elements led us to believe that tRNA recognition by the other archaeal leucyl-tRNAs would be similar to recognition by the halobacterial synthetase. Consequently, these synthetases can be useful in the construction of orthogonal

30   tRNA-synthetase pairs when combined with suppressor tRNAs derived from archaeal leucyl-tRNAs.

[0204]　　　　Because archaeal leucyl-tRNAs and synthetases are highly homologous to one another, both species-specific and cross-species combinations could potentially function as efficient orthogonal tRNA-synthetase pairs. Therefore, each of the five potential orthogonal tRNAs (AfL3, HhL4, MjL2, PfL5, and PhL2) was examined in the presence of

5　each of the six separate archaeal synthetases (AfLRS, ApLRS, HhLRS, MjLRS, MtLRS, and PhLRS) for the ability to suppress A184TAG in *bla*. All five orthogonal tRNAs afforded a higher level of amber suppression in the absence of an archaeal synthetase than is observed when no amber suppressor tRNA is present in the cell. All five suppressors are, therefore, expressed, processed, and functionally charged to some degree by an endogenous

10　*E. coli* synthetase. Nevertheless, only three of the five tRNAs (PhL2, AfL3, and HhL4) gave a higher level of suppression when a foreign synthetase (either MjLRS, MtLRS, or AfLRS) was coexpressed with the tRNA than was observed with no synthetase. The MjL2 and PfL5 suppressors fail to give an enhancement in suppression when coexpressed with a cognate or noncognate archaeal synthetase. Without being limited to one theory, these

15　tRNAs may be expressed as functional suppressor tRNAs in *E. coli* but are unable to be charged due to incompatibility with both cognate and noncognate synthetases. In the case of MjL2, the suppressor is derived from the natural substrate for MjLRS, so it seems unlikely that the tRNA would not be charged, when other tRNAs are efficiently charged by MjLRS. Another explanation might be that the tRNAs are efficiently charged but are

20　incompatible with the *E. coli* translational machinery, but this is not consistent with the fact that some suppression is observed when no archaeal synthetase is present. Another possibility is that MjL2 and PfL5 are efficiently charged with leucine, but are deacylated in an editing process by an endogenous *E. coli* synthetase. In any case, not all archaeal leucyl isoacceptors are equivalent in their ability to function as orthogonal amber suppressors in *E.*

25　*coli*.

[0205]　　　　Only three of the six leucyl-tRNA synthetases (MjLRS, MtLRS, and AfLRS) cloned from archaea gave a higher level of suppression when combined with any of the five orthogonal tRNAs. In the case of HhLRS, the synthetase does not yield protein when over-expressed. Without being limited to one theory, it is most likely, PhLRS and ApLRS do not

30　express functional protein in *E. coli* either, but it is also possible that the proteins are not active at 37°C, or do not recognize any of the orthogonal tRNAs tested. There was no evidence that some tRNAs are preferred substrates for a specific synthetase. Indeed,

although a tRNA from *M. jannaschii* was one of the five orthogonal tRNAs examined, the halobacterium-derived suppressor was the preferred substrate for MjLRS. All three functional tRNAs gave the highest level of suppression when charged by MtLRS or MjLRS, and to a lesser degree with AfLRS.

5 **[0206]**   ·Although on the whole the archaeal leucyl synthetases have similar tRNA recognition properties, it is clear from *in vitro* charging experiments that there are some differences in their recognition of tRNA. The charging of crude total *E. coli* tRNA by AfLRS and MtLRS is only 5- and 13-fold higher, respectively, than the background reaction observed with no synthetase, whereas MjLRS is able to charge *E. coli* tRNA 50-fold over

10 background. Such differences in tRNA recognition among highly homologous synthetases was unanticipated, but not without precedent (Kwok and Wong (1980) Can. J. Biochem. 58, 213-218). Since aminoacyl-tRNA synthetases have evolved only to be orthogonal to the non-cognate tRNAs present in their own host's cytoplasm, it is perhaps not surprising that subtle variations in sequence or chemical modification can lead to mischarging in foreign

15 systems.

[0207]   *Improving the activity of orthogonal suppressor tRNAs.* To date, we have identified and characterized three orthogonal tRNA-synthetase pairs: the yeast glutamine (Liu and Schultz (1999) Proc. Natl. Acad. Sci. U. S. A. 96, 4780-4785), yeast aspartate (Pastrnak et al., (2000) Helv. Chim. Acta 83, 2277-2286), and archaeal tyrosine pairs (Wang

20 et al., (2000) J. Am. Chem. Soc. 122, 5010-5011). Of these systems, only the tyrosine system gives levels of amber suppression comparable to the levels observed for strong native amber suppressors such as supD or supF. When expressed with a the high-copy β-lactamase reporter pBLAM (the reporter plasmid for this study was a medium-copy plasmid) in the presence of their cognate synthetase, cells containing the original glutamine,

25 aspartate, and tyrosine orthogonal amber suppressor tRNAs can survive to 140, 60, and 1220 ug/mL ampicillin, respectively (Pastrnak et al., (2000) Helv. Chim. Acta 83, 2277-2286; Wang et al., (2000) J. Am. Chem. Soc. 122, 5010-5011). A high level of suppression may be critical to the successful modification of the amino acid specificity of synthetases using a double-sieve selection strategy (Liu and Schultz (1999) Proc. Natl. Acad. Sci. U. S.

30 A. 96, 4780-4785). For suppression systems with low activity, it is often difficult to distinguish active and inactive synthetases in selection experiments due to their similarity in phenotype. A high level of suppression is required for the production of protein containing

unnatural amino acids. Therefore, a great deal of attention has been paid to those features
of orthogonal tRNAs that give rise to robust suppression.

[0208] Previous work on frameshift and amber suppression in *E. coli* clearly indicates that
positions 31, 32, 37, and 38 of the tRNA anticodon loop have profound effects on
5   suppression efficiency (Yarus et al., (1986) J. Mol. Biol. 192, 235-255; Smith et al., (1987)
Nucleic Acids Res. 15, 4669-4686; Raftery and Yarus (1987) EMBO J. 6, 1499-1506;
Kleina (1990) J. Mol. Biol. 213, 705-717). The presence of G37 in all the archaeal leucyl
tRNAs led us to believe that a substitution at this position might lead to a higher
suppression efficiency. Indeed, randomization of the anticodon loop showed that the most
10   efficient suppressors have the anticodon loop CUCUAAA. Although the tRNA was toxic,
the G37A mutant also emerged through selection with the *M. jannaschii* tyrosine system
(Wang and Schultz (2001) Chem. Biol. 8, 883-890) as the most potent suppressor thus far
observed for this system. Similar selection experiments with the yeast-derived glutamine
and aspartate orthogonal pairs have been performed in which libraries of positions 32-38 of
15   the anticodon loop are replaced with degenerate bases then subjected to positive ampicillin
selection in the presence of the cognate synthetase (J.C.A., P.G.S., and Miro Pastrnak,
unpublished results). In both cases, the anticodon loop sequence CUCUAAA afforded the
highest suppression efficiency corresponding to six-fold and five-fold enhancements in the
concentration of ampicillin at which growth is observed for the glutamine and aspartate
20   systems, respectively. In at least three other systems, tRNAs with the anticodon loop
sequence CUCUAAA have emerged as the most efficient amber suppressors. The
anticodon loop sequence CUUCCUAA was found to be the most efficient sequence for a
leucyl AGGA suppressor. Selection experiments on tRNAs with randomized anticodon
loops in *E. coli* tRNA$_2^{Ser}$ similarly converged on the anticodon loop sequence CUUCCUAA
25   for AGGA suppression (Magliery et al., (2001) J. Mol. Biol. 307, 755-769), and the
sequence CUUCAAA also emerged as the most efficient anticodon loop sequence for a
leucyl opal suppressor. These results suggest that the preferred anticodon loop sequence is
determined by interactions with endogenous translational machinery rather than the
particular preferences of the aminoacyl-tRNA synthetases. Indeed, the anticodon loop may
30   require sequence-specific modifications in order to function optimally (Soderberg and
Poulter (2000) Biochemistry 39, 6546-6553; Sussman and Kim (1976) Science 192, 853-
858). Alternatively, Yarus (Yarus (1982) Science 218, 646-652) has suggested that the

entire anticodon stem and loop (positions 27-43 of the tRNA) together function as an "extended anticodon" that interacts with ribosome as a module. The entire sequence of this region can help to define the identity of the anticodon for proper decoding.

[0209]   All three codons examined in this study were most efficiently suppressed by tRNAs with the sequence CU(X)XXXAA in the anticodon loop. Although this consensus sequence is preferred for TAG, TGA, and AGGA codons, other sequences may be preferable for other four- and five-base codons. In previous studies (Magliery et al., (2001) J. Mol. Biol. 307, 755-769), the most efficient suppressor tRNAs had bases at positions 32, 33, 37, and 38 which differed from the consensus sequence. For example, the most efficient suppressors of the codon CUAG had an anticodon loop with the sequence CGCTAGGA, deviating at both U33 and A37. In addition, some synthetases employ position 37 as a strong positive determinant for recognition, in which case a CU(X)XXXAA anticodon loop sequence can prove to be non-optimal.

[0210]   Optimization of the anticodon loop sequence as described above was sufficient to provide an efficient amber suppressor tRNA for the leucine system. Optimization of the anticodon loop of the AGGA frameshift suppressors derived from HhL4 also afforded a viable tRNA. However the suppression efficiency (4.6%) of this tRNA, HL(AGGA)1, is far lower than that measured for the suppression of amber codons by HL(TAG)2. Indeed, this suppressor permitted survival at only 75μg/mL ampicillin, significantly less than the seryl AGGA suppressor (Ser2AGGA) identified previously (Magliery et al., (2001) J. Mol. Biol. 307, 755-769), which can survive to 275 ug/mL ampicillin when expressed in plasmid pACKO-A184AGGA. In general, the best AGGA suppressors are less active than the best amber suppressors (Anderson et al., (2002) Chem. Biol. 9, 237-244), but there appears to be something particular to HhL4 that hinders its ability to act as a frameshift suppressor. The only feature obviously different from robust four-base suppressors previously identified (Atkins et al., (1991) Annu. Rev. Genet. 25, 201-228) is the presence of a very large D loop in HhL4. Most suppressors have 9 nucleotides in the D loop and 4 base pairs in the stem. HhL4 has only 3 base pairs in the stem and 13 bases in the loop. Moreover, previous studies have shown the D loop to play a role in frameshift suppression (Tuohy et al., (1992) J. Mol. Biol. 228, 1042-1054). Not only did we see no increase in activity upon randomization of the D loop, there was also a great deal of sequence variation among the most active suppressors.

[0211] The serendipitous appearance of mutations in the G49:U65 base pair of the four-base suppressor tRNAs suggested that non-canonical base pairing in the stem regions of tRNAs has a deleterious effect on suppression efficiency. This hypothesis was further supported by a randomization and selection experiment on the acceptor stem of the HhL4-

5   derived amber suppressor. The three terminal base pairs of the acceptor stem were simultaneously randomized. This library of tRNAs would therefore contain all combinations of mismatched and Watson-Crick base pairs. In fact, 98.4% of the theoretical members of this library should have at least one mismatched base pair. Nevertheless, in the 9 active acceptor stem mutants outlined in **Figure 4, Panel B**, all positions are occupied by

10   Watson-Crick base pairs. Similarly, the D, TψC, anticodon, and acceptor stems of the yeast glutamine amber suppressor tRNA have been individually randomized and subjected to positive selection (J.C.A. and P.G.S., unpublished results). In all surviving clones, every position in these stem regions was occupied by a Watson-Crick pair. In the parent tRNA, the 6:67 base pair is U:G. Mutation of this base pair to U:A results in a doubling of the

15   concentration of ampicillin at which cells can grow. Also, when subjected to positive selection, the only mutations that emerged from random mutagenesis of the leucyl consensus-derived frameshift suppressor appeared at mispaired sites. Others have also noted that mispairing in stem regions adversely affects suppression efficiency (Buttcher et al., (1994) Biochem. Biophys. Res. Commun. 200, 370-377; Hou et al., (1992)

20   Biochemistry 31, 4157-4160). Without being limited to one theory, it may be that tRNAs with mispaired bases are not readily folded into the correct cloverleaf structure and therefore are not readily processed and modified (Furdon et al., (1983) Nucleic Acids Res. 11, 1491-1505). A quantitative analysis of the ratio of charged to uncharged species and of the ratio of fully processed to unprocessed tRNA present in the cell could enhance our

25   understanding of the mechanisms by which these poorly-suppressing tRNAs are impaired.
[0212]   An analysis of multiple sequence alignments of many families of tRNAs reveal multiple examples of conserved non-Watson-Crick pairings. For example, a G3:U70 base pair is a conserved positive determinant for recognition by *E. coli* alanyl-tRNA synthetase (Martinis and Schimmel (1995) in tRNA: Structure, Biosynthesis, and Function

30   (Soll, D., and RajBhandary, U., Eds.) pp 349-370, ASM Press, Washington, DC.). If the element is a conserved positive determinant for recognition, then it may prove difficult to construct robust suppressor tRNAs for the cognate synthetase. Most often, however, the

mispairing present in native sequences is only found in specific isoacceptors. Without being limited to one theory, these and other variations from the consensus sequence of the family of tRNAs present in individual isoacceptors may be present as a result of subtle, species-specific adaptations in positive or negative synthetase recognition, optimal

5   processing and modification, or interactions with elongation factors. Alternatively, these variations may simply be the result of neutral evolutionary drift.

[0213]   When transferred to another species, these variations are unlikely to offer to the new host's translational machinery any advantages they conferred to the source organism. Furthermore, for cross-species pairs, the synthetase is unlikely to recognize any

10   species-specific identity elements present in the tRNA. Only those recognition elements common to the entire family are likely to be useful. Similarly, any processing or modification adaptations particular to a specific tRNA would be lost to the *E. coli* translational apparatus. These variations may even be deleterious to suppression efficiency, particularly when these variations are mismatched bases in stem regions. Suppressors

15   tRNAs derived from the consensus sequence preserve only those features that are broadly shared by the entire family, and eliminate potentially deleterious variations. Therefore, suppressor tRNAs derived from the consensus sequence may in general lead to higher suppression efficiencies.

[0214]   Although optimization of the anticodon loop and elimination of mispairing

20   gave modest to large increases in suppression efficiency, these modifications were not sufficient to provide robust AGGA and opal suppressor tRNAs. Only the consensus-derived suppressors had activities comparable to the $tRNA_2^{Ser}$-derived suppressors described previously (Anderson et al., (2002) Chem. Biol. 9, 237-244). A comparison of the consensus-derived sequences for HL(AGGA)3 and HL(AGGA)2 reveal that there are 14

25   base substitutions, but neither sequence has mispairs. Without being limited to one theory, perhaps by using the consensus sequence of the entire family of tRNAs, those bases that are specific to any particular tRNA and may be detrimental to activity are identified and eliminated.

[0215] *Improving the selectivity of orthogonal suppressor tRNAs.* Unfortunately,

30   improvements in the activity of these suppressor tRNAs also brought about an undesirable increase in the level of suppression observed in the absence of synthetase. The original *M.*

jannaschii tyrosine orthogonal suppressor tRNA was partially charged by an *E. coli* synthetase, but the reaction was eliminated by mutagenesis (Wang and Schultz (2001) Chem. Biol. 8, 883-890). A double sieve selection was able to identify mutants of the wild-type tRNA with excellent orthogonality, but there was also a significant loss of overall

5 activity. Ideally, mutations could be introduced into the tRNA that would eliminate the cross-reactivity with *E. coli* synthetases but preserve high levels of suppression efficiency. Since aminoacyl-tRNA synthetases frequently recognize positions within the acceptor stem and discriminator base of tRNAs (Martinis and Schimmel (1995) in tRNA: Structure, Biosynthesis, and Function (Soll and RajBhandary, Eds.) pp 349-370, ASM Press,

10 Washington, DC.), it is likely that an *E. coli* synthetase that charges the orthogonal tRNA would have a positive recognition element in this region. If this determinant could be changed without destroying recognition by the foreign synthetase, activity could be preserved while eliminating the background reaction. When this strategy was applied to the HhL4-derived amber suppressor, such mutants were indeed found. Several mutants

15 preserved or even improved suppression efficiency when coexpressed with MtLRS but had nearly background levels of amber suppression (7.5 versus 5 ug/mL ampicillin) in the absence of the synthetase. These mutants had reversed the third base pair from G:C to C:G, and an inspection of the recognition elements known for various *E. coli* synthetases suggests the identity of the *E. coli* synthetase that had cross-reacted with HhL4-derived

20 suppressor. Both GlnRS and LysRS of *E. coli* conserve G3:C70 and frequently cross-react with amber suppressor tRNAs (Kleina (1990) J. Mol. Biol. 213, 705-717). Because LysRS also conserves A73, this is the more likely candidate for the cross-reactive *E. coli* synthetase (Freist and Gauss (1995) Biol. Chem. Hoppe-Seyler 376, 451-472; McClain et al., (1988) Science 242, 1681-1684). This strategy can be a general solution to the problem of

25 improving the specificity of cross-reactive orthogonal tRNAs since most *E. coli* aminoacyl-tRNA synthetases contain positive determinants within the acceptor stem.

[0216]   We have shown that the leucyl-tRNA synthetase from the archaean *Methanobacterium thermoautotrophicum* and mutants of a halobacterial tRNA function as an orthogonal pair in *E. coli*. Mutagenesis experiments showed that the two most

30 significant criteria that lead to efficient orthogonal amber suppressor tRNAs are a CU(X)XXXAA anticodon loop and the lack of non-canonical or mismatched base pairs in the stem regions. From these selections, we have identified efficient amber, four-base, and opal orthogonal suppressor tRNAs. We have also devised a consensus strategy to rationally

design efficient orthogonal tRNAs. This leucyl-orthogonal pair can be combined with the *M. jannaschii* pair to site-specifically incorporate two unique unnatural amino acids simultaneously into proteins *in vivo*.

[0217]   *Abbreviations and Textual Footnotes*: Af, Ap, Hh, Mj, Mt , Pf, Ph, and Ec: *Archaeoglobus fulgidus, Aeuropyrum pernix, Halobacterium sp. NRC-1, Methanococcus jannaschii, Methanobacterium thermoautotrophicum, Pyrococcus furiosus, Pyrococcus horikoshi,* and *Escherichia coli*, respectively; LRS, leucyl-tRNA synthetase; *bla*, gene for β-lactamase; *lacZ*, gene for β-galactosidase.

EXAMPLE 2: EXEMPLARY LEUCYL O-RSs AND LEUCYL O-tRNAs.

[0218]   Exemplary O-tRNAs comprise, e.g., SEQ ID NO.:1-7 and 12 (See, Table 3). Exemplary O-RSs include, e.g., SEQ ID NOs.: 15 and 16 (*See*, Table 3). Exemplary polynucleotides that encode O-RSs or portions thereof include, e.g., SEQ ID NOs.: 13 and 14.

[0219]   Further details of the invention, and in particular experimental details, can be found in Anderson, John Christopher, "Pathway Engineering of the Expanding Genetic Code," Ph.D. Dissertation, The Scripps Research Institute [2003].

[0220]   It is understood that the examples and embodiments described herein are for illustrative purposes only and that various modifications or changes in light thereof will be suggested to persons skilled in the art and are to be included within the spirit and purview of this application and scope of the appended claims.

[0221]   While the foregoing invention has been described in some detail for purposes of clarity and understanding, it will be clear to one skilled in the art from a reading of this disclosure that various changes in form and detail can be made without departing from the true scope of the invention. For example, all the techniques and apparatus described above can be used in various combinations. All publications, patents, patent applications, and/or other documents cited in this application are incorporated by reference in their entirety for all purposes to the same extent as if each individual publication, patent, patent application, and/or other document were individually indicated to be incorporated by reference for all purposes.

TABLE 3

SEQ ID:	Label	SEQUENCE
SEQ ID: 1	HL(TAG)1 tRNA	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACT**CTA**GATCCG TTCTCGTAGGAGTTCGAGGGTTCGAATCCCTTCCCTCGCACCA
SEQ ID: 2	HL(TAG)2 tRNA	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACTCTA**A**ATCCG TTCTCGTAGGAGTTCGAGGGTTCGAATCCCTTCCCTCGCACCA
SEQ ID: 3	HL(TAG)3 tRNA	**CCC**AGGGTAGCCAAGCTCGGCCAACGGCGACGGACTCTAAATCCG TTCTCGTAGGAGTTCGAGGGTTCGAATCCCTTCCCT**GGG**ACCA
SEQ ID: 4	HL(AGGA)1 tRNA	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACT**TCCTA**ATCC GTTCTCGTAGGAGTTCGAGGGTTCGAATCCCTTCCCTCGCACCA
SEQ ID: 5	HL(AGGA)2 tRNA	GCGAGGGTAGCCAAGCTCGGCCAACGGCGACGGACTTCCTAATCC GTTCTCGTAGGAGTTCGAGGGTTCGAATCCCT**C**CCCTCGCACCA
SEQ ID: 6	HL(AGGA)3 tRNA	GCGGGGGGTTGCCGAGCCTGGCCAAAGGCGCCGGACTTCCTAATCC **G**GTCCCGTAGGGGTTC**C**GGGGTTCAAATCCCCGCCCCCGCACCA
SEQ ID: 7	HL(TGA)1 tRNA	GCGGGGGGTTGCCGAGCCTGGCCAAAGGCGCCGGACT**TCA**AATCCG GTCCCGTAGGGGTTCCGGGGTTCAAATCCCCGCCCCCGCACCA
SEQ ID: 8	J17 M. jannaschii mtRNA$_{CUA}^{Tyr}$	CCGGCGGTAGTTCAGCAGGGCAGAACGGCGGACTCTAAATCCGCA TGGCGCTGGTTCAAATCCGGCCCGCCGGACCA
SEQ ID: 9	SupD	GGAGAGATGCCGGAGCGGCTGAACGGACCGGTCTCTAAAACCGGA GTAGGGGCAACTCTACCGGGGGTTCAAATCCCCCTCTCTCCGCCA
SEQ ID: 10	Ser2AGGA	GGAGAGATGCCGGAGCGGCTGAACGGACCGGTCTTCCTAAACCGG AGTAGGGGCAACTCTACCGGGGGTTCAAATCCCCCTCTCTCCGCC A
SEQ ID: 11	Leu4 of Halobacterium sp. NRC-1	GCGAGGGUAGCCAAGCUCGGCCAACGGCGACGGACUCAAGAUCCG UUCUCGUAGGAGUUCGAGGGUUCGAAUCCCUUCCCUCGCACCA
SEQ ID: 12	Concensus-derived AGGA Suppressor	GCGGGGGGUUGCCGAGCCUGGCCAAAGGCGCCGGACUUCCUAAUCC CGUCCCGUAGGGGUUCGGGGGUUCAAAUCCCCGCCCCCGCACCA
SEQ ID: 13	Archaeoglobus fulgidus leucyl tRNA-synthetase (AFLRS)	ATGAGCGATT TCAGGATAAT TGAGGAGAAG TGGCAGAAGG CGTGGGAGAA GGACAGAATT TTTGAGTCCG ATCCTAATGA GAAGGAGAAG TTTTTTCTCA CAATTCCCTA TCCTTACCTT AATGGAAATC TTCACGCAGG TCACACGAGA ACCTTCACAA TTGGCGATGC CTTCGCCAGA TACATGAGAA TGAAGGGCTA CAACGTTCTC TTTCCCCTCG GCTTTCATGT TACGGGCACC CCAATCATTG GCCTTGCGGA GCTCATAGCC AAGAGGGACG AGAGGACGAT AGAGGTTTAC ACCAAATACC ATGACGTTCC GCTGGAGGAC TTGCTTCAGC TCACAACTCC AGAGAAAATC GTTGAGTACT TCTCAAGGGA GGCGCTGCAG GCTTTGAAGA GCATAGGCTA CTCCATTGAC TGGAGGAGGG TTTTCACCAC AACCGATGAA

		GAGTATCAGA GATTCATCGA GTGGCAGTAC TGGAAGCTCA AGGAGCTTGG
		CCTGATTGTG AAGGGCACCC ACCCCGTCAG ATACTGCCCC CACGACCAGA
		ATCCTGTTGA AGACCACGAC CTTCTCGCTG GGGAGGAGGC AACTATTGTT
		GAATTTACCG TTATAAAGTT CAGGCTTGAA GATGGAGACC TCATTTTCCC
		CTGTGCAACT CTCCGTCCCG AAACCGTGTT TGGCGTCACG AACATCTGGG
		TAAAGCCGAC AACCTACGTA ATTGCCGAGG TGGATGGGGA AAAGTGGTTT
		GTGAGCAAAG AGGCTTACGA GAAGCTCACC TACACGGAGA AAAAAGTCAG
		GCTGCTGGAG GAGGTTGATG CGTCGCAGTT CTTCGGCAAG TACGTCATAG
		TCCCGCTGGT AAACAGAAAA GTGCCAATTC TGCCTGCAGA GTTTGTTGAC
		ACCGACAACG CAACAGGAGT TGTGATGAGC GTTCCCGCAC ACGCTCCTTT
		TGACCTGGCT GCCATTGAGG ACTTGAAGAG AGACGAGGAA ACGCTGGCGA
		AGTACGGAAT TGACAAAAGC GTTGTAGAGA GCATAAAGCC AATAGTTCTG
		ATTAAGACGG ACATTGAAGG TGTTCCTGCT GAGAAGCTAA TAAGAGAGCT
		TGGAGTGAAG AGCCAGAAGG ACAAGGAGCT GCTGGATAAG GCAACCAAGA
		CCCTCTACAA GAAGGAGTAC CACACGGGAA TCATGCTGGA CAACACGATG
		AACTATGCTG GAATGAAAGT TTCTGAGGCG AAGGAGAGAG TTCATGAGGA
		TTTGGTTAAG CTTGGCTTGG GGGATGTTTT CTACGAGTTC AGCGAGAAGC
		CCGTAATCTG CAGGTGCGGA ACGAAGTGCG TTGTTAAGGT TGTTAGGGAC
		CAGTGGTTCC TGAACTACTC CAACAGAGAG TGGAAGGAGA AGGTTCTGAA
		TCACCTTGAA AAGATGCGAA TCATCCCCGA CTACTACAAG GAGGAGTTCA
		GGAACAAGAT TGAGTGGCTC AGGGACAAGG CTTGTGCCAG AAGGAAGGGG
		CTTGGAACGA GAATTCCGTG GGATAAGGAG TGGCTCATCG AGAGCCTTTC
		AGACTCAACA ATCTACATGG CCTACTACAT CCTTGCCAAG TACATCAACG
		CAGGATTGCT CAAGGCCGAG AACATGACTC CCGAGTTCCT CGACTACGTG
		CTGCTGGGCA AAGGTGAGGT TGGGAAAGTT GCGGAAGCTT CAAAACTCAG
		CGTGGAGTTA ATCCAGCAGA TCAGGGACGA CTTCGAGTAC TGGTATCCCG
		TTGACCTAAG AAGCAGTGGC AAGGACTTGG TTGCAAACCA CCTGCTCTTC
		TACCTCTTCC ACCACGTCGC CATTTTCCCG CCAGATAAGT GGCCGAGGGC
		AATTGCCGTA AACGGATACG TCAGCCTTGA GGGCAAGAAG ATGAGCAAGA
		GCAAAGGGCC CTTGCTAACG ATGAAGAGGG CGGTGCAGCA GTATGGTGCG
		GATGTGACGA GGCTCTACAT CCTCCACGCT GCAGAGTACG ACAGCGATGC
		GGACTGGAAG AGCAGAGAGG TTGAAGGGCT TGCAAACCAC CTCAGGAGGT
		TCTACAACCT CGTGAAGGAG AACTACCTGA AAGAGGTGGG AGAGCTAACA
		ACCCTCGACC GCTGGCTTGT GAGCAGGATG CAGAGGGCAA TAAAGGAAGT
		GAGGGAGGCT ATGGACAACC TGCAGACGAG GAGGGCCGTG AATGCCGCCT
		TCTTCGAGCT CATGAACGAC GTGAGATGGT ATCTGAGGAG AGGAGGTGAG
		AACCTCGCTA TAATACTGGA CGACTGGATC AAGCTCCTCG CCCCCTTTGC
		TCCGCACATT TGCGAGGAGC TGTGGCACTT GAAGCATGAC AGCTACGTCA
		GCCTCGAAAG CTACCCAGAA TACGACGAAA CCAGGGTTGA CGAGGAGGCG
		GAGAGAATTG AGGAATACCT CCGAAACCTT GTTGAGGACA TTCAGGAAAT
		CAAGAAGTTT GTTAGCGATG CGAAGGAGGT TTACATTGCT CCCGCCGAAG
		ACTGGAAGGT TAAGGCAGCA AAGGTCGTTG CTGAAAGCGG GGATGTTGGG
		GAGGCGATGA AGCAGCTTAT GCAGGACGAG GAGCTTAGGA AGCTCGGCAA
		AGAAGTGTCA AATTTCGTCA AGAAGATTTT CAAAGACAGA AAGAAGCTGA
		TGCTAGTTAA GGAGTGGGAA GTTCTGCAGC AGAACCTGAA ATTTATTGAG
		AATGAGACCG GACTGAAGGT TATTCTTGAT ACTCAGAGAG TTCCTGAGGA
		GAAGAGGAGG CAGGCAGTTC CGGGCAAGCC CGCGATTTAT GTTGCTTAA
SEQ ID: 14	Methanob acterium thermoau totrophi cum leucyl tRNA-syntheta se (MtLRS)	GTGGATATTG AAAGAAAATG GCGTGATAGA TGGAGAGATG CTGGCATATT
		TCAGGCTGAC CCTGATGACA GAGAAAAGAT ATTCCTCACA GTCGCTTACC
		CCTACCCCAG TGGTGCGATG CACATAGGAC ACGGAGGGAC CTACACTGTC
		CCTGATGTCT ATGCACGGTT CAAGAGGATG CAGGGCTACA CGTCCTGTT
		TCCCATGGCC TGGCATGTCA CAGGGGCCCC TGTCATAGGG ATAGCGCGGA
		GGATTCAGAG GAAGGATCCC TGGACCCTCA AAATCTACAG GGAGGTCCAC
		AGGGTCCCCG AGGATGAGCT TGAACGTTTC AGTGACCCTG AGTACATAGT
		TGAATACTTC AGCAGGGAAT ACCGGTCTGT TATGGAGGAT ATGGGCTACT
		CCATCGACTG GAGGCGTGAA TTCAAAACCA CGGATCCCAC CTACAGCAGG
		TTCATACAGT GGCAGATAAG GAAGCTGAGG GACCTTGGCC TCGTAAGGAA
		GGGCGCCCAT CCTGTTAAGT ACTGCCCTGA ATGTGAAAAC CCTGTGGGTG
		ACCATGACCT CCTTGAGGGT GAGGGGGTTG CCATAAACCA GCTCACACTC

| | | | CTCAAATTCA AACTTGGAGA CTCATACCTG GTCGCAGCCA CCTTCAGGCC
CGAGACAATC TATGGGGCCA CCAACCTCTG GCTGAACCCT GATGAGGATT
ATGTGAGGGT TGAAACAGGT GGTGAGGAGT GGATAATAAG CAGGGCTGCC
GTGGATAATC TTTCACACCA GAAACTGGAC CTCAAGGTTT CCGGTGACGT
CAACCCCGGG GACCTGATAG GGATGTGCGT GGAGAATCCT GTGACGGGCC
AGGAACACCC CATACTCCCG GCTTCCTTCG TTGACCCTGA ATATGCCACA
GGTGTTGTGT TCTCTGTCCC TGCACATGCC CCTGCAGACT TCATAGCCCT
TGAGGACCTC AGGACAGACC ATGAACTCCT TGAAAGGTAC GGTCTTGAGG
ATGTGGTTGC TGATATTGAG CCCGTGAATG TCATAGCAGT GGATGGCTAC
GGTGAGTTCC CGGCGGCCGA GGTTATAGAG AAATTTGGTG TCAGAAACCA
GGAGGACCCC CGCCTTGAGG ATGCCACCGG GGAGCTATAC AAGATCGAGC
ATGCGAGGGG TGTTATGAGC AGCCACATCC CTGTCTATGG TGGTATGAAG
GTCTCTGAGG CCCGTGAGGT CATCGCTGAT GAACTGAAGG ACCAGGGCCT
TGCAGATGAG ATGTATGAAT CGCTGAGCG ACCTGTTATA TGCCGCTGCG
GTGGCAGGTG CGTTGTGAGG GTCATGGAGG ACCAGTGGTT CATGAAGTAC
TCTGATGACG CCTGGAAGGA CCTCGCCCAC AGGTGCCTCG ATGGCATGAA
GATAATACCC GAGGAGGTCC GGGCCAACTT TGAATACTAC ATCGACTGGC
TCAATGACTG GGCATGTTCA AGGAGGATAG GCCTTGGAAC AAGGCTGCCC
TGGGATGAGA GGTGGATCAT CGAACCCCTC ACAGACTCAA CAATCTACAT
GGCATATTAC ACCATCGCAC ACCGCCTCAG GGAGATGGAT GCCGGGGAGA
TGGACGATGA GTTCTTTGAT GCCATATTCC TAGATGATTC AGGAACCTTT
GAGGATCTCA GGGAGGAATT CCGGTACTGG TACCCCCTTG ACTGGAGGCT
CTCTGCAAAG GACCTCATAG GCAATCACCT GACATTCCAT ATATTCCACC
ACTCAGCCAT ATTCCCTGAG TCAGGGTGGC CCCGGGGGGC TGTGGTCTTT
GGTATGGGCC TTCTTGAGGG CAACAAGATG TCATCCTCCA AGGGCAACGT
CATACTCCTG AGGGATGCCA TCGAGAAGCA CGGTGCAGAC GTGGTGCGGC
TCTTCCTCAT GTCCTCAGCA GAGCCATGGC AGGACTTTGA CTGGAGGGAG
AGTGAGGTCA TCGGGACCCG CAGGAGGATT GAATGGTTCA GGGAATTCGG
AGAGAGGGTC TCAGGTATCC TGGATGGTAG GCCAGTCCTC AGTGAGGTTA
CTCCAGCTGA ACCTGAAAGC TTCATTGGAA GGTGGATGAT GGGTCAGCTG
AACCAGAGGA TACGTGAAGC CACAAGGGCC CTTGAATCAT TCCAGACAAG
AAAGGCAGTT CAGGAGGCAC TCTATCTCCT TAAAAAGGAT GTTGACCACT
ACCTTAAGCG TGTTGAGGGT AGAGTTGATG ATGAGGTTAA ATCTGTCCTT
GCAAACGTTC TGCACGCCTG GATAAGGCTC ATGGCTCCAT TCATACCCTA
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